

2-PHENYL-1-[4-(2-AMINOETHOXY)-BENZYL-INDOLES  
AS ESTROGENIC AGENTS

This application claims the benefit of U.S. Application No. 60/015,553, filed  
5 April 1996 and is a continuation-in-part of that prior application.

The present invention relates to new 2-Phenyl-1-[4-(2-Aminoethoxy)-Benzyl]-Indole compounds which are useful as estrogenic agents, as well as pharmaceutical compositions and methods of treatment utilizing these compounds.

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Background of the Invention

The use of hormone replacement therapy for bone loss prevention in post-menopausal women is well preceded. The normal protocol calls for estrogen supplementation using such formulations containing estrone, estriol, ethynodiol or conjugated estrogens isolated from natural sources (i.e. Premarin® conjugated estrogens from Wyeth-Ayerst). In some patients, therapy may be contraindicated due to the proliferative effects of unopposed estrogens (estrogens not given in combination with progestins) have on uterine tissue. This proliferation is associated with increased risk for endometriosis and/or endometrial cancer. The effects of unopposed estrogens on breast tissue is less clear, but is of some concern. The need for estrogens which can maintain the bone sparing effect while minimizing the proliferative effects in the uterus and breast is evident. Certain nonsteroidal antiestrogens have been shown to maintain bone mass in the ovariectomized rat model as well as in human clinical trials. Tamoxifen (sold as Novadex® brand tamoxifen citrate by Zeneca Pharmaceuticals, Wilmington, Delaware), for example, is a useful palliative for the treatment of breast cancer and has been demonstrated to exert an estrogen agonist-like effect on the bone, in humans. However, it is also a partial agonist in the uterus and this is cause for some concern. Raloxifene, a benzothiophene antiestrogen, has been shown to stimulate uterine growth in the ovariectomized rat to a lesser extent than Tamoxifen while maintaining the ability to spare bone. A suitable review of tissue selective estrogens is seen in the article "Tissue-Selective Actions Of Estrogen Analogs", *Bone* Vol. 17, No. 4, October 1995, 181S-190S.

The use of indoles as estrogen antagonists has been reported by Von Angerer, Chemical Abstracts, Vol. 99, No. 7 (1983), Abstract No. 53886u. Also, see, J.Med.Chem. 1990, 33, 2635-2640; J.Med.Chem. 1987, 30, 131-136. Also see Ger. 5 Offen., DE 3821148 A1 891228 and WO 96/03375. These prior art compounds share some structural similarities with the present compounds, but are functionally different. For compounds containing a basic amine, there is no phenyl group to rigidify the side chain. The reported data for these compounds indicates that they may have a weaker binding to estrogen receptor than the compounds of the present invention and the 10 reported compounds containing the basic side chain show some uterotrophic effect in the rat uterus. One compound from the listed family of compounds in WO 96/03375 possesses a benzyl group, but does not have a basic side chain. The majority of these compounds fall into a class of compounds best characterized as being "pure antiestrogens". Many of the compounds described presently, due to their particular 15 side chain, act as pure antiestrogens in the uterus, however, show strong estrogenic action in the bone and cardiovascular systems. No such action is demonstrated for the related prior art compounds described herein.

WO A 95 17383 (Kar Bio AB) describes indole antiestrogens with long straight 20 chains. Another related patent WO A 93 10741 describes 5-Hydroxyindole with a generic descriptor incorporating other side chains. WO 93/23374 (Otsuka Pharmaceuticals, Japan) describes compounds which differ from the present invention; where R<sub>3</sub> in the present formulas I and II, below, is defined as thioalkyl and the reference discloses no such compounds having chains from the indole nitrogen having 25 the same structure as the ones provided by the present invention. Where the side chain claimed is similar to that described herein, the compounds are amides: Acylated indoles are not claimed in the present invention.

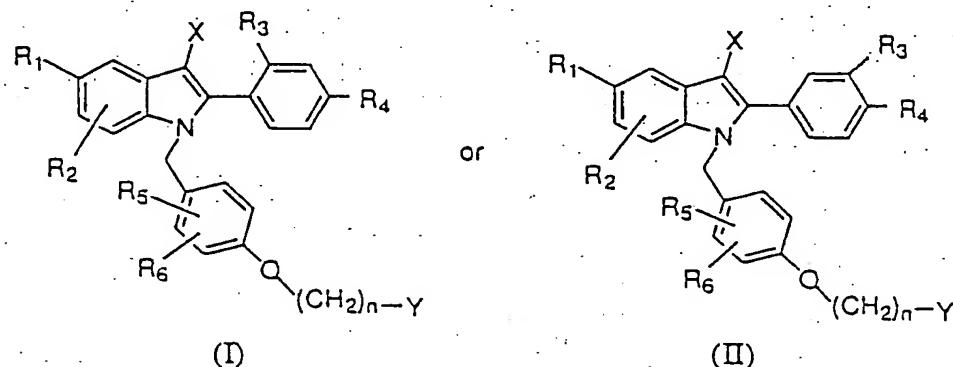
#### Description of the Invention

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2-Phenylindoles of the general structure type shown in formulas (I) and (II) are estrogen agonists/antagonists useful for the treatment of diseases associated with estrogen deficiency. The compounds of the present invention show strong binding to

the estrogen receptor. *In vitro* assays, including an Ishikawa alkaline phosphatase assay and an ERE transfection assay, show these compounds are antiestrogens with little to no intrinsic estrogenicity and they have proven capable of completely antagonizing the effects of 17 $\beta$ -estradiol while showing little or no uterine stimulation in a rat uterine assay when dosed alone. Additionally, some of these compounds are capable of inhibiting bone loss in an ovariectomized rat while showing little or no uterine stimulation. These compounds also decrease the weight gain normally seen in the ovariectomized animals as well as reduce total cholesterol levels.

10 The present invention includes compounds of the formulas I or II, below:



15 wherein:

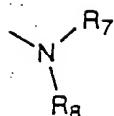
$R_1$  is selected from H, OH or the  $C_1$ - $C_{12}$  esters (straight chain or branched) or  $C_1$ - $C_{12}$  (straight chain or branched or cyclic) alkyl ethers thereof, or halogens; or  $C_1$ - $C_4$  halogenated ethers including trifluoromethyl ether and trichloromethyl ether.

20  $R_2, R_3, R_4, R_5$ , and  $R_6$  are independently selected from H, OH or the C<sub>1</sub>-C<sub>12</sub> esters (straight chain or branched) or C<sub>1</sub>-C<sub>12</sub> alkyl ethers (straight chain or branched or cyclic) thereof, halogens, or C<sub>1</sub>-C<sub>4</sub> halogenated ethers including trifluoromethyl ether and trichloromethyl ether, cyano, C<sub>1</sub>-C<sub>6</sub> alkyl (straight chain or branched), or trifluoromethyl, with the proviso that, when  $R_1$  is H,  $R_2$  is not OH.

25 X is selected from H, C<sub>1</sub>-C<sub>6</sub> alkyl, cyano, nitro, trifluoromethyl, halogen;  
n is 2 or 3.

Y is selected from:

a) the moiety:



wherein R<sub>7</sub> and R<sub>8</sub> are independently selected from the group of H, C<sub>1</sub>-

5 C<sub>6</sub> alkyl, or phenyl optionally substituted by CN, C<sub>1</sub>-C<sub>6</sub> alkyl (straight chain or branched), C<sub>1</sub>-C<sub>6</sub> alkoxy (straight chain or branched), halogen, -OH, -CF<sub>3</sub>, or -OCF<sub>3</sub>;

b) a five-membered saturated, unsaturated or partially unsaturated heterocycle containing up to two heteroatoms selected from the group consisting of -O-,  
10 -NH-, -N(C<sub>1</sub>C<sub>4</sub> alkyl)-, -N=, and -S(O)<sub>m</sub>-, wherein m is an integer of from 0-2,  
optionally substituted with 1-3 substituents independently selected from the group  
consisting of hydrogen, hydroxyl, halo, C<sub>1</sub>-C<sub>4</sub> alkyl, trihalomethyl, C<sub>1</sub>-C<sub>4</sub> alkoxy,  
trihalomethoxy, C<sub>1</sub>-C<sub>4</sub> acyloxy, C<sub>1</sub>-C<sub>4</sub> alkylthio, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub>  
alkylsulfonyl, hydroxy (C<sub>1</sub>-C<sub>4</sub>)alkyl, -CO<sub>2</sub>H-, -CN-, -CONHR<sub>1</sub>-, -NH<sub>2</sub>-, C<sub>1</sub>-C<sub>4</sub>  
15 alkylamino, di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, -NHSO<sub>2</sub>R<sub>1</sub>-, -NHCOR<sub>1</sub>-, -NO<sub>2</sub>, and phenyl  
optionally substituted with 1-3 (C<sub>1</sub>-C<sub>4</sub>)alkyl;

c) a six-membered saturated, unsaturated or partially unsaturated heterocycle containing up to two heteroatoms selected from the group consisting of -O-,  
20 -NH-, -N(C<sub>1</sub>C<sub>4</sub> alkyl)-, -N=, and -S(O)<sub>m</sub>-, wherein m is an integer of from 0-2,  
optionally substituted with 1-3 substituents independently selected from the group  
consisting of hydrogen, hydroxyl, halo, C<sub>1</sub>-C<sub>4</sub> alkyl, trihalomethyl, C<sub>1</sub>-C<sub>4</sub> alkoxy,  
trihalomethoxy, C<sub>1</sub>-C<sub>4</sub> acyloxy, C<sub>1</sub>-C<sub>4</sub> alkylthio, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub>  
alkylsulfonyl, hydroxy (C<sub>1</sub>-C<sub>4</sub>)alkyl, -CO<sub>2</sub>H-, -CN-, -CONHR<sub>1</sub>-, -NH<sub>2</sub>-, C<sub>1</sub>-C<sub>4</sub>  
25 alkylamino, di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, -NHSO<sub>2</sub>R<sub>1</sub>-, -NHCOR<sub>1</sub>-, -NO<sub>2</sub>, and phenyl  
optionally substituted with 1-3 (C<sub>1</sub>-C<sub>4</sub>)alkyl;

d) a seven-membered saturated, unsaturated or partially unsaturated heterocycle containing up to two heteroatoms selected from the group  
30 consisting of -O-, -NH-, -N(C<sub>1</sub>C<sub>4</sub> alkyl)-, -N=, and -S(O)<sub>m</sub>-, wherein m is an integer  
of from 0-2, optionally substituted with 1-3 substituents independently selected from  
the group consisting of hydrogen, hydroxyl, halo, C<sub>1</sub>-C<sub>4</sub> alkyl, trihalomethyl, C<sub>1</sub>-C<sub>4</sub>

alkoxy, trihalomethoxy, C<sub>1</sub>-C<sub>4</sub> acyloxy, C<sub>1</sub>-C<sub>4</sub> alkylthio, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, hydroxy (C<sub>1</sub>-C<sub>4</sub>)alkyl, -CO<sub>2</sub>H-, -CN-, -CONHR<sub>1</sub>-, -NH<sub>2</sub>-, C<sub>1</sub>-C<sub>4</sub> alkylamino, di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, -NHSO<sub>2</sub>R<sub>1</sub>-, -NHCOR<sub>1</sub>-, -NO<sub>2</sub>, and phenyl optionally substituted with 1-3 (C<sub>1</sub>-C<sub>4</sub>)alkyl; or

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e) a bicyclic heterocycle containing from 6-12 carbon atoms either bridged or fused and containing up to two heteroatoms selected from the group consisting of -O-, -NH-, -N(C<sub>1</sub>C<sub>4</sub> alkyl)-, and -S(O)<sub>m</sub>-, wherein m is an integer of from 0-2, optionally substituted with 1-3 substituents independently selected from the group consisting of hydrogen, hydroxyl, halo, C<sub>1</sub>-C<sub>4</sub> alkyl, trihalomethyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, trihalomethoxy, C<sub>1</sub>-C<sub>4</sub> acyloxy, C<sub>1</sub>-C<sub>4</sub> alkylthio, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, hydroxy (C<sub>1</sub>-C<sub>4</sub>)alkyl, -CO<sub>2</sub>H-, -CN-, -CONHR<sub>1</sub>-, -NH<sub>2</sub>-, C<sub>1</sub>-C<sub>4</sub> alkylamino, di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, -NHSO<sub>2</sub>R<sub>1</sub>-, -NHCOR<sub>1</sub>-, -NO<sub>2</sub>, and phenyl optionally substituted with 1-3 (C<sub>1</sub>-C<sub>4</sub>) alkyl;

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and the pharmaceutically acceptable salts thereof.

The more preferred compounds of this invention are those having the general structures I or II, above, wherein:

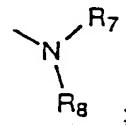
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R<sub>1</sub> is selected from H, OH or the C<sub>1</sub>-C<sub>12</sub> esters or alkyl ethers thereof, halogen; R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub>, R<sub>5</sub>, and R<sub>6</sub> are independently selected from H, OH or the C<sub>1</sub>-C<sub>12</sub> esters or alkyl ethers thereof, halogen, cyano, C<sub>1</sub>-C<sub>6</sub> alkyl, or trihalomethyl, preferably trifluoromethyl, with the proviso that, when R<sub>1</sub> is H, R<sub>2</sub> is not OH;

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X is selected from H, C<sub>1</sub>-C<sub>6</sub> alkyl, cyano, nitro, trifluoromethyl, halogen;

Y is the moiety



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R<sub>7</sub> and R<sub>8</sub> are selected independently from H, C<sub>1</sub>-C<sub>6</sub> alkyl, or combined by -(CH<sub>2</sub>)<sub>p</sub>-, wherein p is an integer of from 2 to 6, so as to form a ring, the ring being optionally substituted by up to three substituents selected from the group of hydrogen, hydroxyl, halo, C<sub>1</sub>-C<sub>4</sub> alkyl, trihalomethyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, trihalomethoxy, C<sub>1</sub>-C<sub>4</sub>

alkylthio, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, hydroxy (C<sub>1</sub>-C<sub>4</sub>)alkyl, -CO<sub>2</sub>H, -CN, -CONH(C<sub>1</sub>-C<sub>4</sub>), -NH<sub>3</sub>, C<sub>1</sub>-C<sub>4</sub> alkylamino, C<sub>1</sub>-C<sub>4</sub> dialkylamino, -NHSO<sub>2</sub>(C<sub>1</sub>-C<sub>4</sub>), -NHCO(C<sub>1</sub>-C<sub>4</sub>), and -NO<sub>3</sub>;

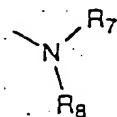
and the pharmaceutically acceptable salts thereof.

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The rings formed by a concatenated R<sub>7</sub> and R<sub>8</sub>, mentioned above, may include, but are not limited to, aziridine, azetidine, pyrrolidine, piperidine, hexamethyleneamine or heptamethyleneamine rings.

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The most preferred compounds of the present invention are those having the structural formulas I or II, above, wherein R<sub>1</sub> is OH; R<sub>2</sub> - R<sub>6</sub> are as defined above; X is selected from the group of Cl, NO<sub>2</sub>, CN, CF<sub>3</sub>, or CH<sub>3</sub>; and Y is the moiety



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and R<sub>7</sub> and R<sub>8</sub> are concatenated together as -(CH<sub>2</sub>)<sub>r</sub>, wherein r is an integer of from 4 to 6, to form a ring optionally substituted by up to three substituents selected from the group of hydrogen, hydroxyl, halo, C<sub>1</sub>-C<sub>4</sub> alkyl, trihalomethyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, trihalomethoxy, C<sub>1</sub>-C<sub>4</sub> alkylthio, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, hydroxy (C<sub>1</sub>-C<sub>4</sub>)alkyl, -CO<sub>2</sub>H, -CN, -CONH(C<sub>1</sub>-C<sub>4</sub>)alkyl, -NH<sub>2</sub>, C<sub>1</sub>-C<sub>4</sub> alkylamino, di(C<sub>1</sub>-C<sub>4</sub>)alkylamino, -NHSO<sub>2</sub>(C<sub>1</sub>-C<sub>4</sub>)alkyl, -NHCO(C<sub>1</sub>-C<sub>4</sub>)alkyl, and -NO<sub>2</sub>;

and the pharmaceutically acceptable salts thereof.

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In another embodiment of this invention, when R<sub>7</sub> and R<sub>8</sub> are concatenated together as -(CH<sub>2</sub>)<sub>p</sub>-, wherein p is an integer of from 2 to 6, preferably 4 to 6, the ring so formed is optionally substituted with 1-3 substituents selected from a group containing C<sub>1</sub>-C<sub>3</sub> alkyl, trifluoromethyl, halogen, hydrogen, phenyl, nitro, -CN.

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The invention includes sulfate, sulfonates and sulfate esters of phenolic groups. Sulfates can be readily prepared by the reaction of the free phenolic compounds with sulfur trioxide complexed with an amine such as pyridine, trimethylamine, triethylamine, etc. Sulfonates can be prepared by treating the free phenolic compound

with the desired amino or alkylamino or dialkylamino sulfamyl chloride in the presence of a suitable base such as pyridine. Sulfate esters can be prepared by reaction of the free phenol with the desired alkanesulfonyl chloride in the presence of a suitable base such as pyridine. Additionally, this invention includes compounds containing phosphates at the phenol as well as dialkyl phosphates. Phosphates can be prepared by reaction of the phenol with the appropriate chlorophosphate. The dialkylphosphates can be hydrolyzed to yield the free phosphates. Phosphinates are also claimed where the phenol is reacted with the desired dialkylphosphinic chloride to yield the desired dialkylphosphinate of the phenol.

10

The invention includes acceptable salt forms formed from the addition reaction with either inorganic or organic acids. Inorganic acids such as hydrochloric acid, hydrobromic acid, hydroiodic acid, sulfuric acid, phosphoric acid, nitric acid useful as well as organic acids such as acetic acid, propionic acid, citric acid, maleic acid, malic acid, tartaric acid, phthalic acid, succinic acid, methanesulfonic acid, toluenesulfonic acid, naphthalenesulfonic acid, camphorsulfonic acid, benzenesulfonic acid are useful. It is known that compounds possessing a basic nitrogen can be complexed with many different acids (both protic and non-protic) and usually it is preferred to administer a compound of this invention in the form of an acid addition salt. Additionally, this invention includes quaternary ammonium salts of the compounds herein. These can be prepared by reacting the nucleophilic amines of the side chain with a suitably reactive alkylating agent such as an alkyl halide or benzyl halide.

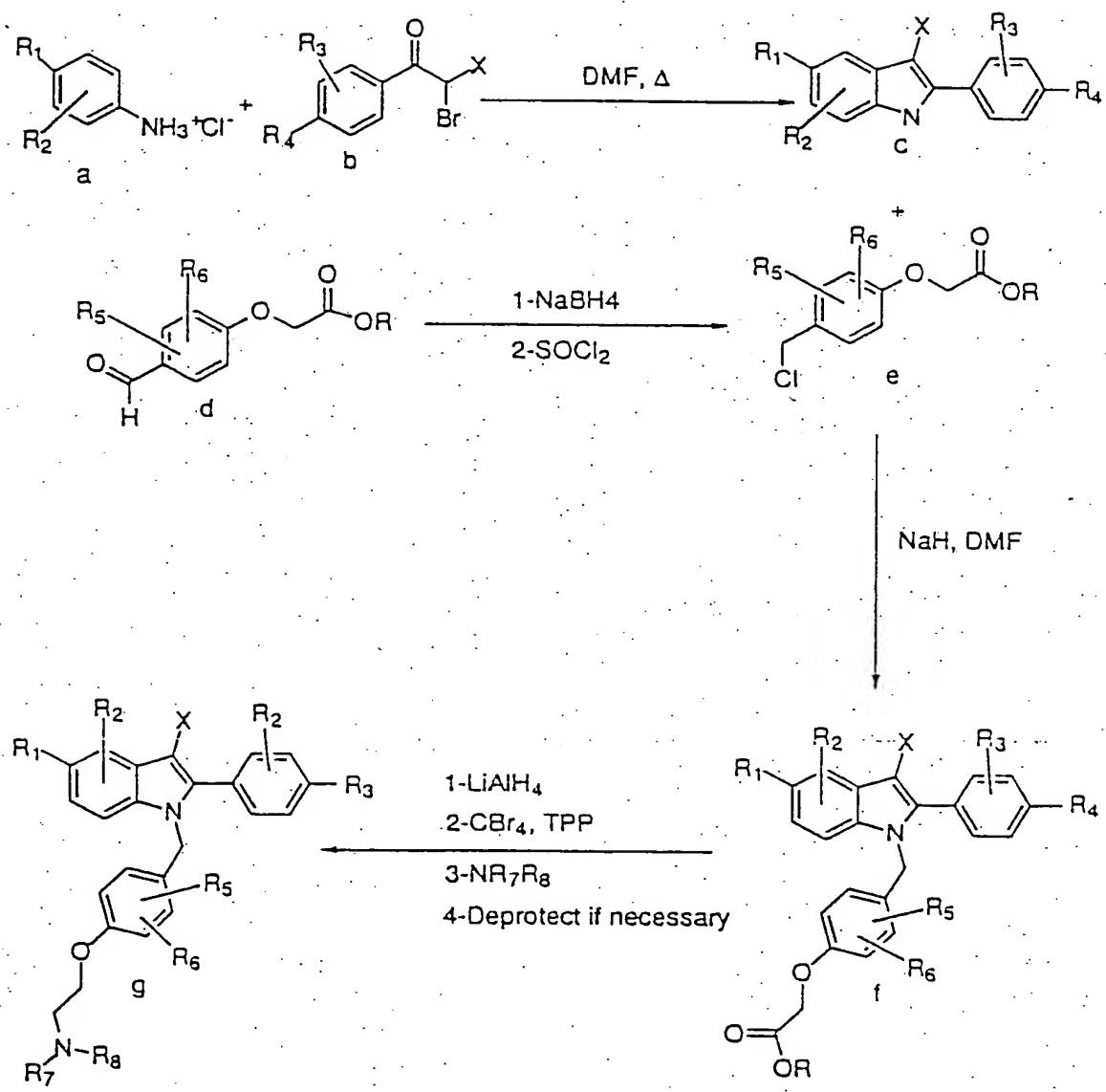
### Methods

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Compounds of this invention can be synthesized in a general sense according to Scheme 1, below.

- 8 -

Scheme 1



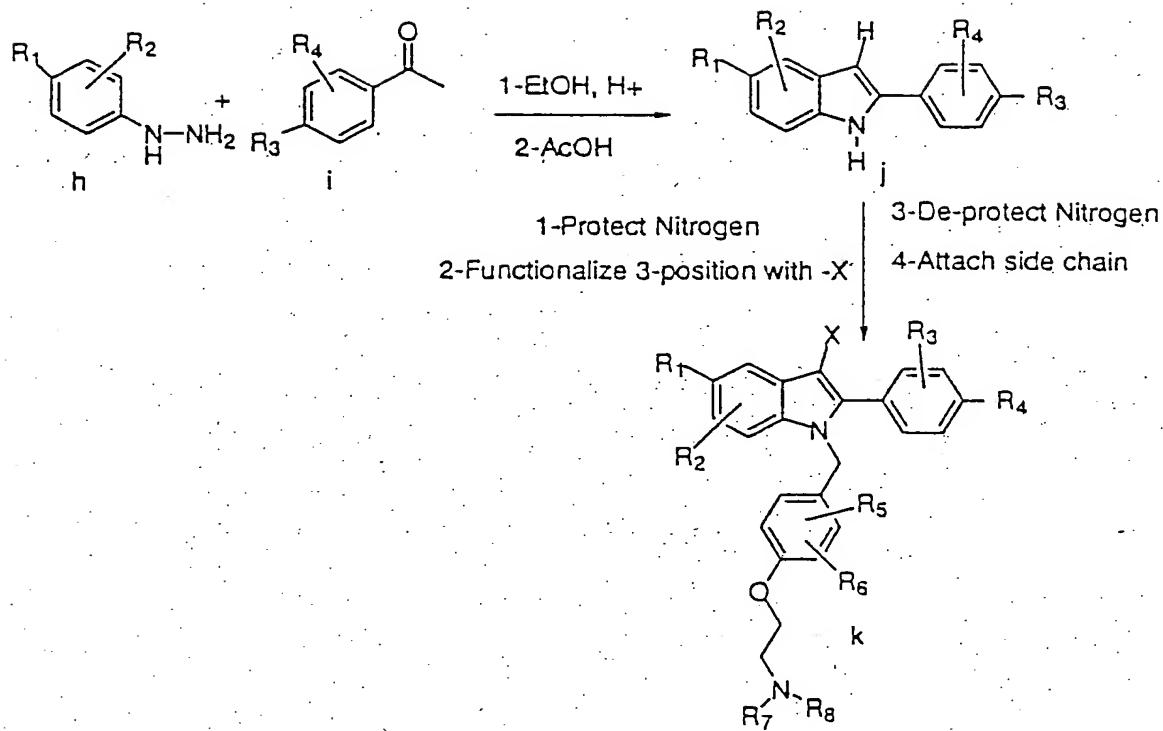
5. The initial indole synthesis is accomplished by heating an appropriately substituted alpha-bromo ketone (b) with the desired aniline (a) in DMF to form the indole (c). The product is then alkylated with a benzyl chloride (e) to give the substituted indole (f). The benzyl chloride (e) can be readily prepared from the aldehyde (d) in 2 steps as given. Product (g) can be prepared from (f) by reduction of

the ester, conversion of the alcohol to a bromide, displacement of the bromide with the desired amine in a suitable solvent such as THF or DMF, and finally, deprotection if necessary. Deprotection is necessary when either R<sub>1</sub> or R<sub>2</sub> or both is a protected phenol. The preferred protecting group is a benzyl group which can be conveniently 5 removed by several conventional methods, especially hydrogenolysis.

For the synthesis of compounds with X=H, halogen, trifluoromethyl, cyano, nitro, an alternative synthesis shown in scheme 2 may be preferable. The formation of halogens at the 3-position can be easily performed with such reagents as N-chlorosuccinamide, N-bromosuccinamide, or N-iodosuccinamide. A 3-Iodoindole 10 compound obtained can be used as a precursor to the 3-trifluoromethyl compound by a coupling reaction utilizing a palladium catalyst and bis(trifluoromethyl) mercury (II). A compound with a cyano group in the 3-position can be prepared by electrophilic cyanation or alternatively the 3-position can be formylated (with a formyl iminium salt, for example) then the formyl group converted to an oxime and subsequently dehydrated 15 to a nitrile. Alternatively, the 3-cyano compound can be synthesized by reaction of the 3-unsubstituted indole with chlorosulfonylisocyanate followed by triethylamine. A compound with the nitro group in the 3-position can be prepared by treating the indole with sodium nitrite and acetic acid. One skilled in the art recognizes these routes are not limiting and other routes are also available.

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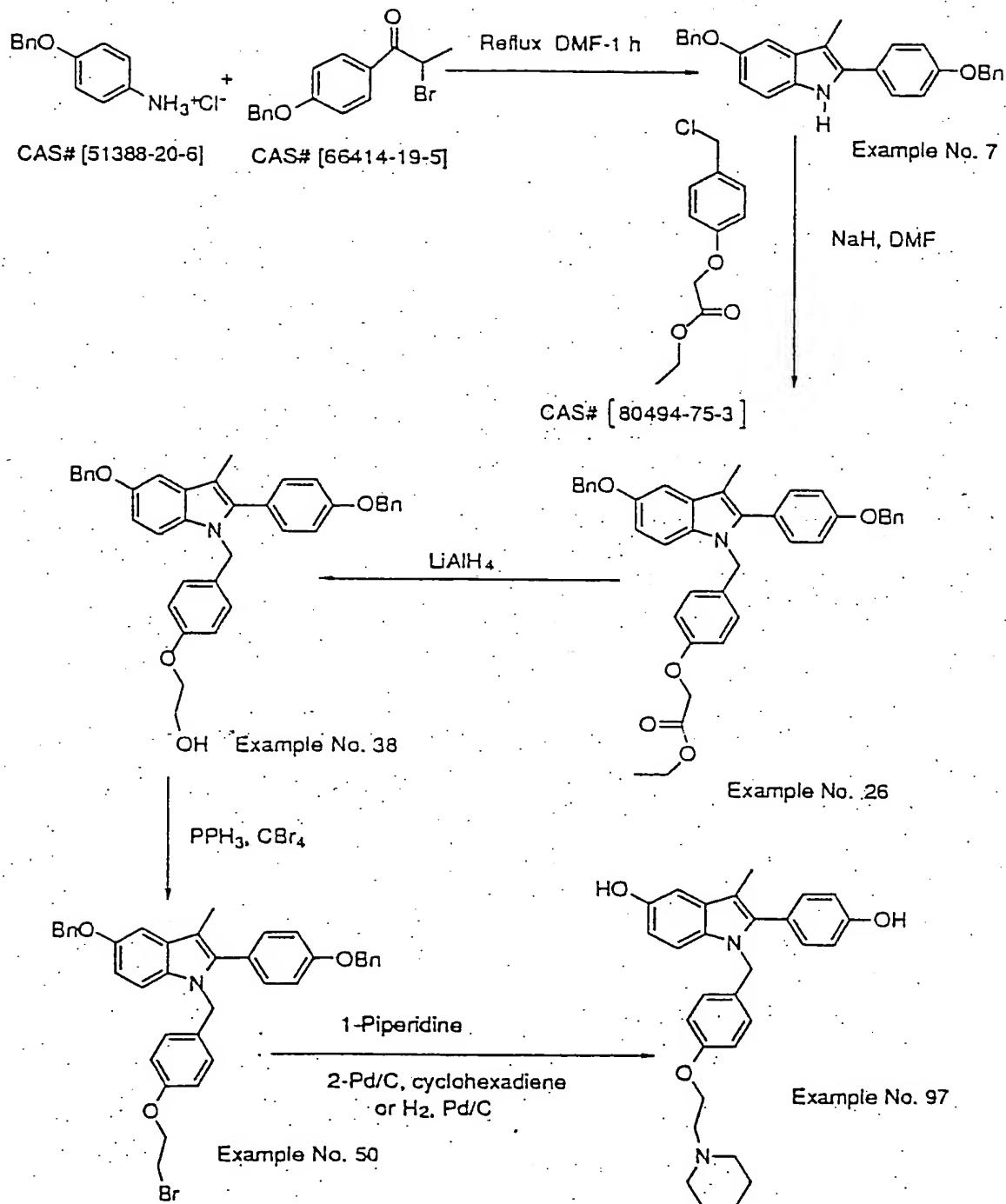
Scheme 2



5      Synthesis of selected representative examples are given in the following schemes:

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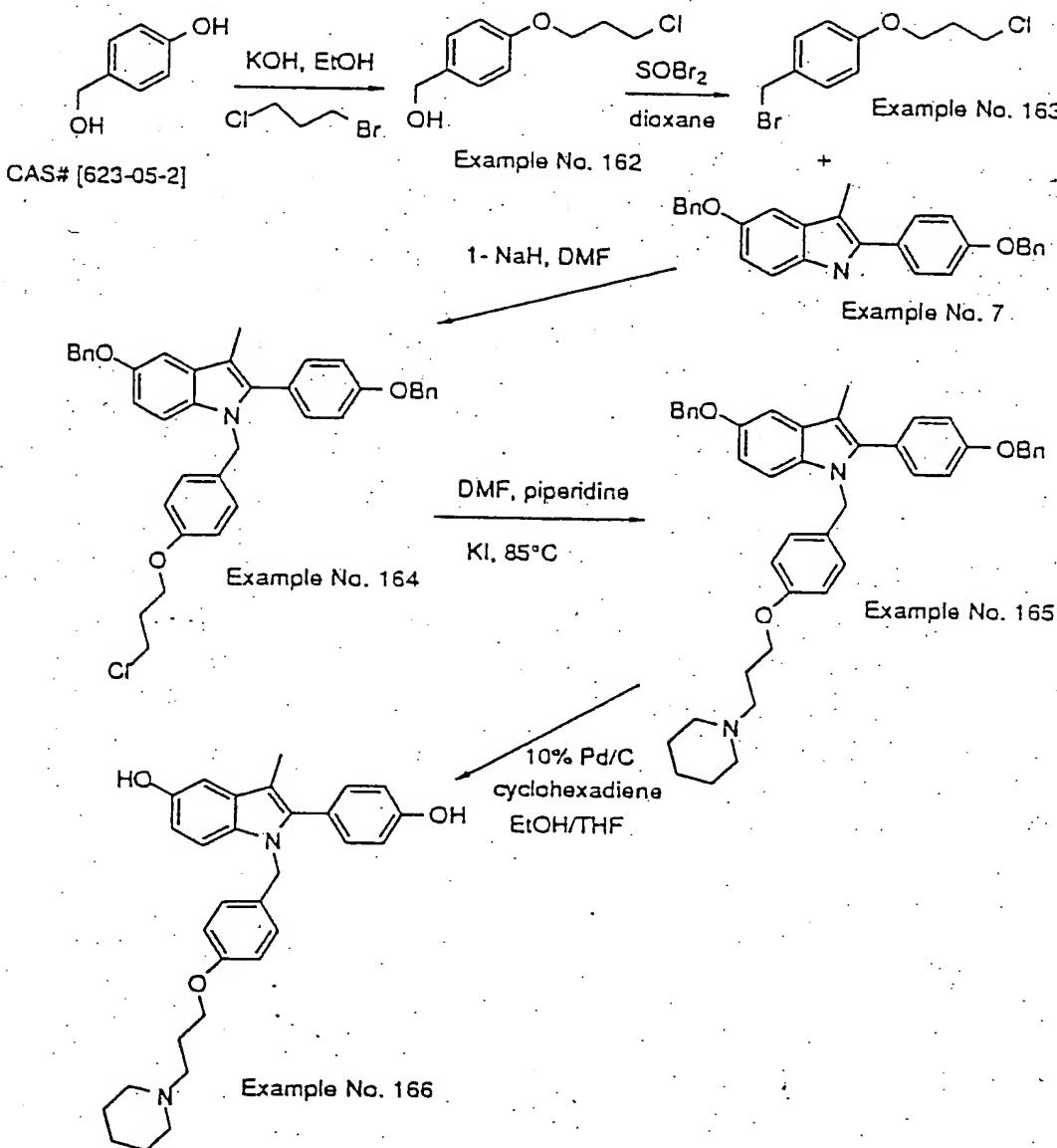
Scheme 3



The synthesis of analogues with a 3-carbon chain (example No. 166) between the oxygen and the basic amine can be accomplished as shown in scheme 4.

Scheme 4

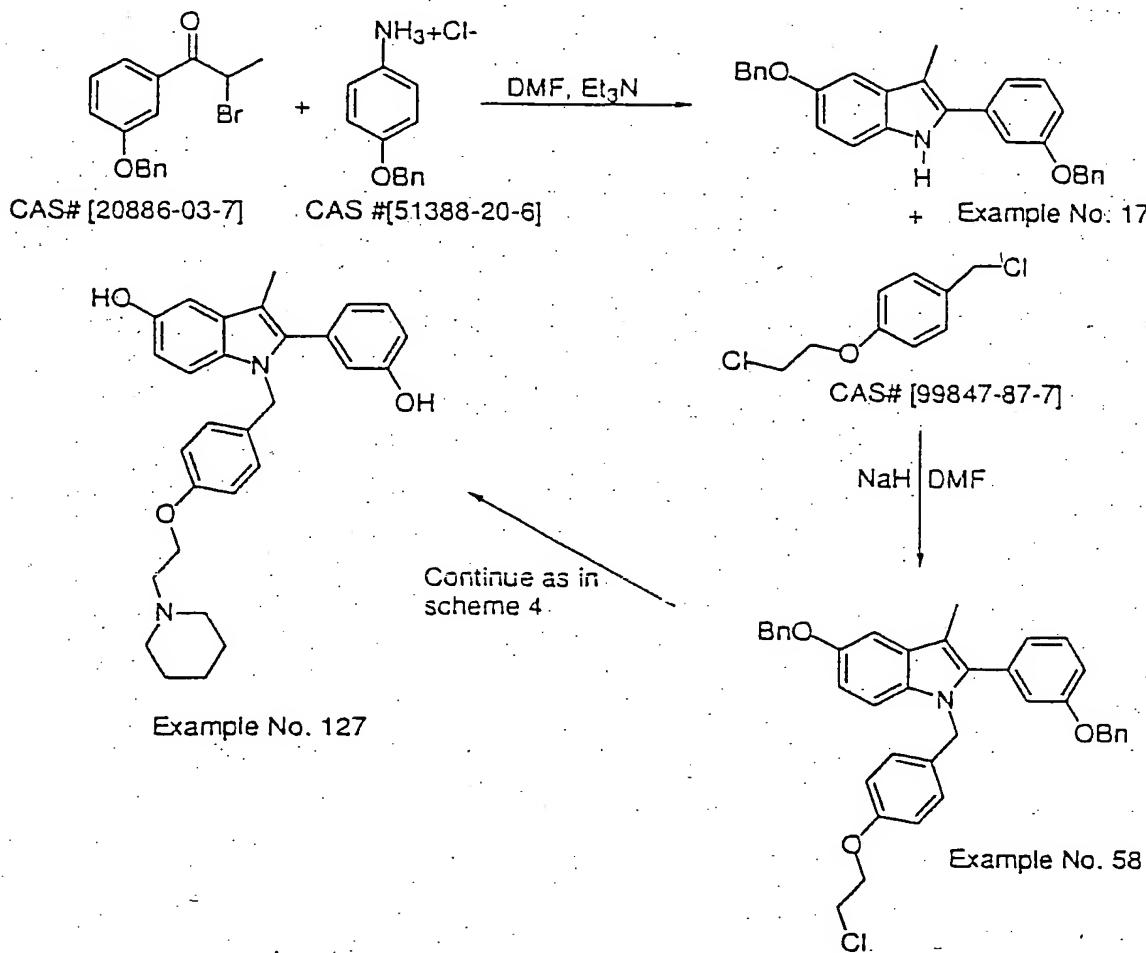
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The synthetic procedure shown in scheme 4 may be used for compounds with two carbon chains analogous to example No. 97 in scheme 3. This is shown in scheme 4a for the synthesis of example No. 127.

5 Scheme 4a

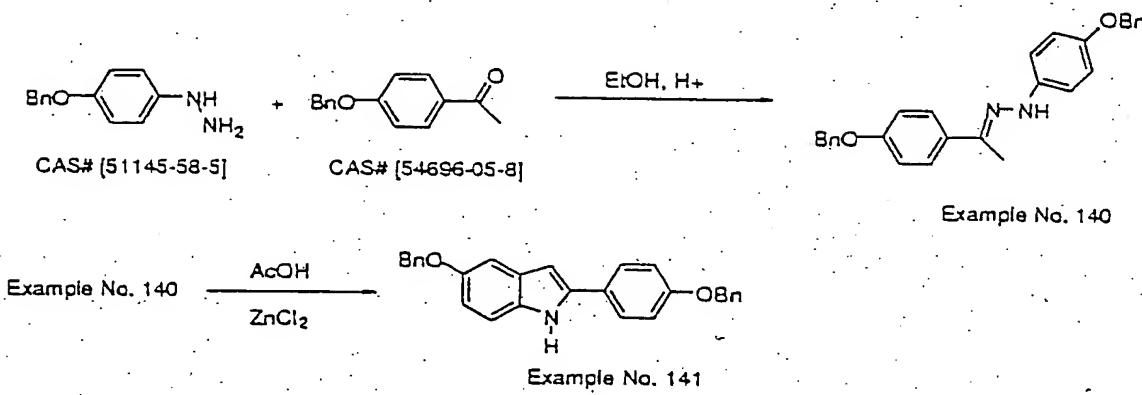


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The synthesis of indoles with alternative substituents (CN, Cl) at the 3-position of the indole both utilize the 3-unsubstituted indole No. 141 for a precursor. The indole is synthesized by the Fisher method utilizing the hydrazone derived from the condensation of 4-benzyloxyacetophenone CAS No. [54696-05-8] and 4-benzyloxyphenylhydrazine

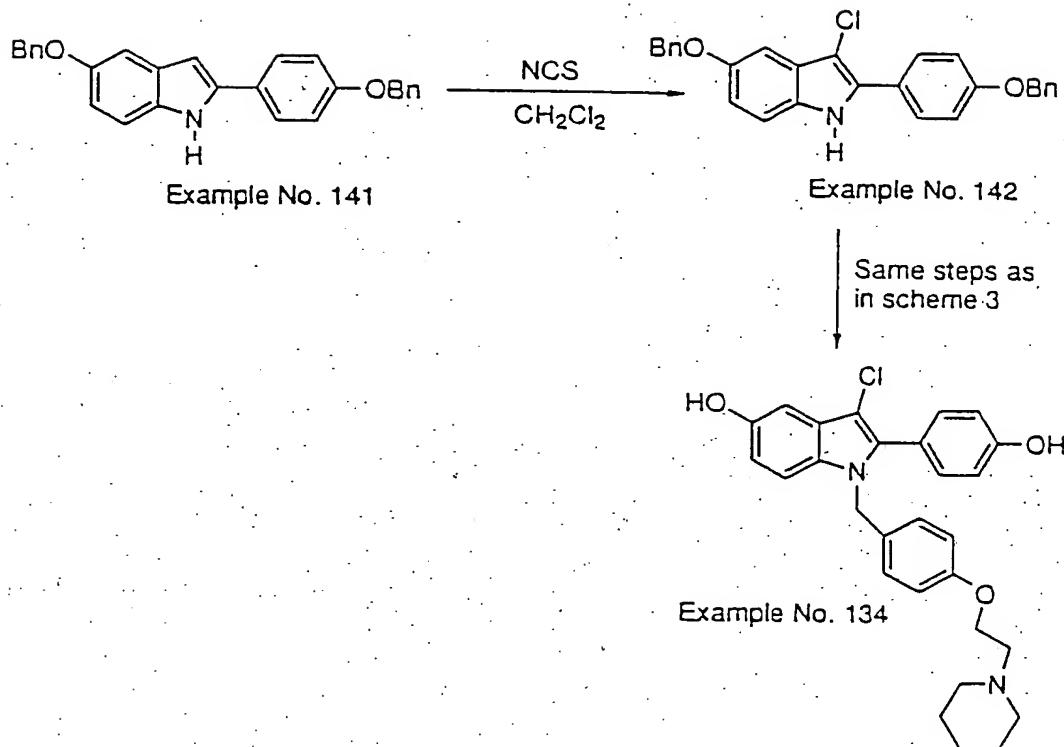
CAS No. [51145-58-5]. The hydrazone No. 140 is then cyclized in acetic acid using zinc chloride to afford the desired indole No. 141. This synthesis can be seen in scheme 5.

5 Scheme 5



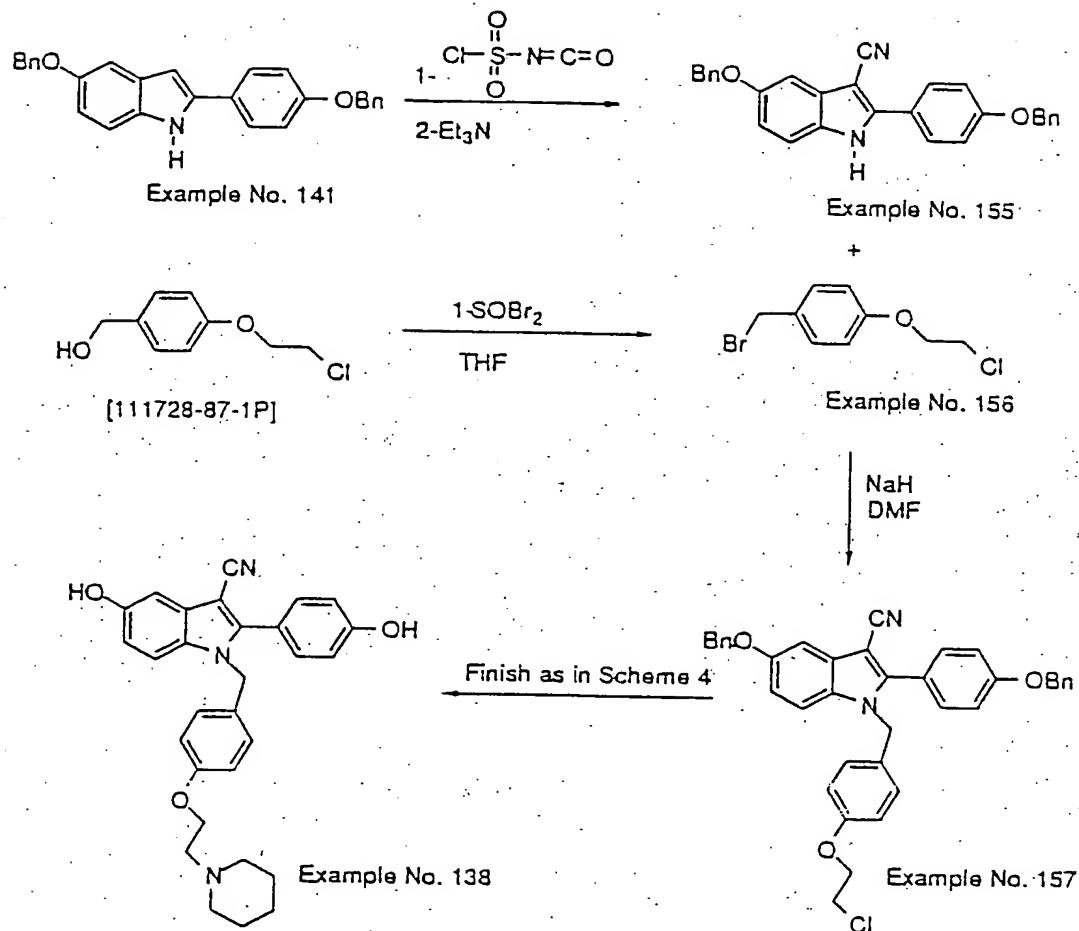
The synthesis of 3-Chloroindole compounds is demonstrated for example No. 134 and shown, *infra*, in scheme 6. The indole No. 141 from scheme 5 is chlorinated with N-chlorosuccinamide. The 3-Chloroindole No. 142, thus obtained, is taken to the final product in analogous fashion to that shown in scheme 3.

Scheme 6



3-Cyano analogues are synthesized from the precursor indole No. 141 as shown in Scheme 7. Reaction of the precursor indole No. 141 with chlorosulfonyl isocyanate followed by addition of triethylamine yields the 3-Cyanoindole No. 155. The side chain is made by conversion of the benzylic alcohol of CAS No. [111728-87-1] to the benzylic bromide No. 156 using thionyl bromide in THF. The indole is alkylated by the side chain in DMF using sodium hydride to give the intermediate No. 157. This can then be taken to the final product No. 138 in an analogous fashion to that shown in scheme 4.

Scheme 7



5 The compounds of the invention are selective estrogen agonists and display high affinity for the estrogen receptor. Unlike many estrogens, however, many of these compounds do not cause increases in uterine wet weight. These compounds are antiestrogenic in the uterus and can completely antagonize the trophic effects of estrogen agonists in uterine tissue. Due to the tissue selective nature of these 10 compounds, they are useful in treating or preventing in a mammal disease states or syndromes which are caused or associated with an estrogen deficiency (in certain tissues such as bone or cardiovascular) or an excess of estrogen (in the uterus or mammary glands). They may also be used in methods of treatment for diseases or

disorders which result from proliferation or abnormal development, actions or growth of endometrial or endometrial-like tissues.

The present compounds have the ability to behave like estrogen agonists by 5 lowering cholesterol and preventing bone loss. These compounds are useful for treating many maladies which result from estrogen effects and estrogen excess or deficiency including osteoporosis, prostatic hypertrophy, male pattern baldness, vaginal and skin atrophy, acne, dysfunctional uterine bleeding, endometrial polyps, benign breast disease, uterine leiomyomas, adenomyosis, ovarian cancer, infertility, 10 breast cancer, endometriosis, endometrial cancer, polycystic ovary syndrome, cardiovascular disease, contraception, Alzheimer's disease, cognitive decline and other CNS disorders, as well as certain cancers including melanoma, prostate cancer, cancers of the colon, CNS cancers, among others. Additionally, these compounds can be used for contraception in pre-menopausal women, as well as hormone replacement 15 therapy in post-menopausal women or in other estrogen deficiency states where estrogen supplementation would be beneficial. They may also be used in disease states where amenorrhea is advantageous, such as leukemia, endometrial ablations, chronic renal or hepatic disease or coagulation diseases or disorders.

20 The compounds of this invention may also be used in methods of treatment for and prevention of bone loss, which may result from an imbalance in a individual's formation of new bone tissues and the resorption of older tissues, leading to a net loss of bone. Such bone depletion results in a range of individuals, particularly in post-menopausal women, women who have undergone bilateral oophorectomy, those 25 receiving or who have received extended corticosteroid therapies, those experiencing gonadal dysgenesis, and those suffering from Cushing's syndrome. Special needs for bone, including teeth and oral bone, replacement can also be addressed using these compounds in individuals with bone fractures, defective bone structures, and those receiving bone-related surgeries and/or the implantation of prosthesis. In addition to 30 those problems described above, these compounds can be used in treatments for osteoarthritis, hypocalcemia, hypercalcemia, Paget's disease, osteomalacia, osteohalisteresis, multiple myeloma and other forms of cancer having deleterious effects on bone tissues. Methods of treating the maladies listed herein are understood

to comprise administering to an individual in need of such treatment a pharmaceutically effective amount of one or more of the compounds of this invention or a pharmaceutically acceptable salt thereof. This invention also includes pharmaceutical compositions utilizing one or more of the present compounds, and/or the 5 pharmaceutically acceptable salts thereof, along with one or more pharmaceutically acceptable carriers, excipients, etc.

It is understood that the dosage, regimen and mode of administration of these 10 compounds will vary according to the malady and the individual being treated and will be subject to the judgement of the medical practitioner involved. It is preferred that the administration of one or more of the compounds herein begin at a low dose and be increased until the desired effects are achieved.

Effective administration of these compounds may be given at a dose of from 15 about 0.1 mg/day to about 1,000 mg/day. Preferably, administration will be from about 10 mg/day to about 600 mg/day, more preferably from about 50 mg/day to about 600 mg/day, in a single dose or in two or more divided doses. Such doses may be administered in any manner useful in directing the active compounds herein to the recipient's bloodstream, including orally, via implants, parenterally (including 20 intravenous, intraperitoneal and subcutaneous injections), rectally, vaginally, and transdermally. For the purposes of this disclosure, transdermal administrations are understood to include all administrations across the surface of the body and the inner linings of bodily passages including epithelial and mucosal tissues. Such 25 administrations may be carried out using the present compounds, or pharmaceutically acceptable salts thereof, in lotions, creams, foams, patches, suspensions, solutions, and suppositories (rectal and vaginal).

Oral formulations containing the active compounds of this invention may 30 comprise any conventionally used oral forms, including tablets, capsules, buccal forms, troches, lozenges and oral liquids, suspensions or solutions. Capsules may contain mixtures of the active compound(s) with inert fillers and/or diluents such as the pharmaceutically acceptable starches (e.g. corn, potato or tapioca starch), sugars, artificial sweetening agents, powdered celluloses, such as crystalline and

microcrystalline celluloses, flours, gelatins, gums, etc. Useful tablet formulations may be made by conventional compression, wet granulation or dry granulation methods and utilize pharmaceutically acceptable diluents, binding agents, lubricants, disintegrants, suspending or stabilizing agents, including, but not limited to, magnesium stearate, 5 stearic acid, talc, sodium lauryl sulfate, microcrystalline cellulose, carboxymethylcellulose calcium, polyvinylpyrrolidone, gelatin, alginic acid, acacia gum, xanthan gum, sodium citrate, complex silicates, calcium carbonate, glycine, dextrin, sucrose, sorbitol, dicalcium phosphate, calcium sulfate, lactose, kaolin, mannitol, sodium chloride, talc, dry starches and powdered sugar. Oral formulations 10 herein may utilize standard delay or time release formulations to alter the absorption of the active compound(s). Suppository formulations may be made from traditional materials, including cocoa butter, with or without the addition of waxes to alter the suppository's melting point, and glycerin. Water soluble suppository bases, such as polyethylene glycols of various molecular weights, may also be used.

15

Aldrich Sure Seal<sup>TM</sup> Solvents, anhydrous without further purification, may be used for the reactions described herein and may be obtained from Aldrich Chemical Company. All reactions were carried out under a nitrogen atmosphere. Chromatography was performed using 230-400 mesh silica gel (Merck Grade 60, 20 Aldrich Chemical Company). Thin layer chromatography was performed with Silica Gel 60 F254 plates from EM Science. <sup>1</sup>H NMR spectra were obtained on a Bruker AM-400 or Bruker DPX-300 instrument in DMSO and chemical shifts reported in ppm. Melting points were determined on a Thomas-Hoover apparatus and are uncorrected. IR spectra were recorded on a Perkin-Elmer diffraction grating or Perkin-Elmer 784 25 spectrophotometers. Mass spectra were recorded on a Kratos MS 50 or Finnigan 8230 mass spectrometers. Elemental analyses were obtained with a Perkin-Elmer 2400 elemental analyzer. Compounds for which CHN are reported are within 0.4% of the theoretical value for the formula given unless expressed otherwise. Compound nomenclature was generally arrived at by use of the Beilstein Autonom<sup>TM</sup> program.

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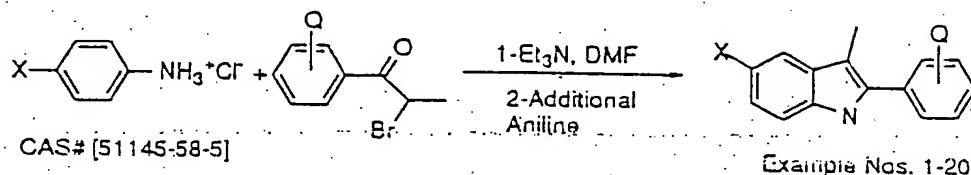
Synthesis of  $\alpha$ -bromo ketones

Method a

The synthesis of the alpha bromo ketones is conveniently accomplished by  
 5 simply dissolving the starting phenyl ketone in ethyl ether (0.05-0.10 M) and at room temperature, 1.1 equivalents of bromine is added in dropwise. The reaction can be monitored by TLC for consumption of starting materials. The reaction is worked up by washing with an aqueous sodium bicarbonate solution followed by a 10% aqueous sodium sulfite solution. The ether layer is washed with brine and dried over  
 10 magnesium sulfate. Concentration of the reaction mixture typically yields the bromoketones in good yield and purity. The bromoketones were taken "as is" (without purification or characterization) to the next step.

3-Methyl indoles

15 Scheme 8



20

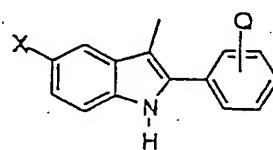


Table 1

Example No.	X	Q
No. 1	H	H
No. 1a	F	OBn
No. 2	H	4'-OBn

Table 1 (Cont'd)

Example No.	X	Q
No. 3	OBn	H
No. 4	OBn	4'-OMe
No. 5	OMe	4'-OMe
No. 6	OBn	4'-OEt
No. 7	OBn	4'-OBn
No. 8	OBn	4'-F
No. 9	OBn	3'-OMe,4'-OBn
No. 10	OBn	3',4'-OCH <sub>2</sub> O-
No. 11	OBn	4'-O-iPr
No. 12	OBn	4'-O-Cp
No. 13	OBn	4'-CF <sub>3</sub>
No. 14	OBn	4'-CH <sub>3</sub>
No. 15	OBn	4'-Cl
No. 16	OBn	2'-OMe,4'-OMe
No. 17	OBn	3'-OBn
No. 18	OBn	4'-OBn,3'-F
No. 19	OBn	3'-OMe
No. 20	OBn	4'-OCF <sub>3</sub>

Method 1

5

Illustrated For Example No. 7

5-Benzylxy-2-(4-benzylxy-phenyl)-3-methyl-1H-indole

A flask was charged with 4-benzylxyaniline hydrochloride CAS No. [51145-58-5]. (45 g, 0.23 mol), 4'-benzylxy-2-bromophenylpropiophenone CAS No. [66414-19-5] (21g, 0.066 mol), and 50 mL DMF. The reaction was heated at reflux for 30 minutes and then cooled to rt and then partitioned between 250 mL EtOAc and 100 mL 1N HCl (aq). The EtOAc was washed with NaHCO<sub>3</sub> (aq) and brine, then dried over MgSO<sub>4</sub>. The solution was concentrated and the residue taken up in CH<sub>2</sub>Cl<sub>2</sub>

and hexanes added to precipitate out 25g of a crude solid. The solid was dissolved in CH<sub>2</sub>Cl<sub>2</sub> and evaporated onto silica gel and chromatographed using CH<sub>2</sub>Cl<sub>2</sub>/Hexane (1:5) to yield 9.2 g of a tan solid (33%): Mp = 150-152°C; <sup>1</sup>H NMR (DMSO) 10.88 (s, 1 H), 7.56 (d, 2 H, J = 8.8 Hz), 7.48 (d, 4 H, J = 7.9 Hz), 7.42-7.29 (m, 6 H), 7.21 (d, 1 H, J = 7.0 Hz), 7.13 (d, 2 H, J = 8.8 Hz), 7.08 (d, 1 H, J = 2.2 Hz), 6.94 (dd, 1 H, J = 8.8, 2.4 Hz), 5.16 (s, 2 H), 5.11 (s, 2 H), 2.33 (s, 3 H); IR (KBr) 3470, 2880, 2820, 1620 cm<sup>-1</sup>; MS eI m/z 419.

10

Method 2 (shown in scheme 8)

Also Illustrated For Example No. 7

Reagents used were same as in method 1 except the additional use of triethylamine in this method. The bromoketone CAS No. [66414-19-5] (50.0 g, 0.16 mol) in 200 mL DMF was treated with the aniline hydrochloride CAS No. [51145-58-5] (44 g, 0.22 mol) and the reaction purged with nitrogen for about 10 minutes. The triethylamine (54.6 mL) was added and the reaction was heated at 120°C for 2 hours. TLC analysis (EtOAc/hexanes) shows the starting material has dissappeared forming a more polar spot. The reaction mixture is allowed to cool down and an additional 48 g. of the aniline hydrochloride was added. The reaction was heated to 150°C for 2 hours. An additional 5 grams of the aniline hydrochloride was added and the reaction was heated at 150°C for an additional 30 minutes. The reaction mixture is allowed to cool to room temperature and then poured into approximately 1.5 liters of water and extracted with 2 liters of ethyl acetate. Solids are dissolved with additional ethyl acetate as neccessary. The ethyl acetate layer is washed with 1 liter of 1 N NaOH solution aq., 1 liter of water, brine, then dried over magnesium sulfate and filtered. The organic layers were concentrated down to yield a crude solid which is stirred with 500 mL of methanol and filtered. This solid is then stirred with 500 mL of ethyl ether and filtered. The solid is stirred alternatively with methanol and ether until it is of whitish color and has a melting point similar to that described for No. 7 in method 1. Reaction yields 36 grams of product.

### Physical Data for Indoles

The following 3-methyl indoles (No. 1- No. 20) were synthesized according to the procedure outlined in scheme 2 using method 2 using the appropriately substituted 5 bromoketones (prepared as given above) and anilines (commercially available; Aldrich) as starting materials.

#### Example No. 1 2-Phenyl-3-methyl-1H-indole

Mp = 90 - 94°C;  $^1\text{H}$  NMR (DMSO) 11.13 (s, 1 H), 7.68 - 7.64 (m, 2 H), 7.54 - 7.46 (m, 3 H), 7.37 - 7.32 (m, 2 H), 7.12 - 7.06 (m, 1 H), 7.03 - 6.97 (m, 1 H), 2.40 (s, 3 H); MS eI m/z 207 (M $^+$ ).

#### Example No. 1a 5-Fluoro-2-(4-benzylxy-phenyl)-3-methyl-1H-indole

Mp = 143 - 146°C.

15

#### Example No. 2 2-(4-Benzylxy-phenyl)-3-methyl-1H-indole

Mp = 118 - 120°C;  $^1\text{H}$  NMR (DMSO) 11.03 (s, 1 H), 7.57 (dd, 2 H, J = 2.0 Hz, 6.6 Hz), 7.48 - 7.46 (m, 3 H), 7.44 - 7.28 (m, 4 H), 7.18 - 7.11 (m, 2 H), 7.08 - 7.03 (m, 1 H), 7.0 - 6.95 (m, 1 H), 5.16 (s, 2 H), 2.36 (s, 3 H); MS eI m/z 313 (M $^+$ ).

#### Example No. 3 5-Benzylxy-2-phenyl-3-methyl-1H-indole

Mp = 141-144°C;  $^1\text{H}$  NMR (DMSO) 10.98 (s, 1 H), 7.65-7.61 (m, 2 H), 7.51-7.44 (m, 4 H), 7.42-7.28 (m, 4 H), 7.23 (d, 1 H, J = 8.8 Hz), 7.10 (d, 1 H, J = 2.5 Hz), 6.80 (d, 1 H, J = 6.0 Hz), 5.10 (s, 2 H), 2.36 (s, 3 H); MS eI m/z 313 (M $^+$ ).

#### Example No. 4 5-Benzylxy-2-(4-methoxy-phenyl)-3-methyl-1H-indole

Mp = 158°C;  $^1\text{H}$  NMR 10.85 (brs, 1 H), 7.56 (d, 2 H, J = 8.8 Hz), 7.48 (d, 2 H, J = 8.3 Hz), 7.45 - 7.36 (m, 2 H), 7.34 - 7.28 (m, 1 H), 7.21 (d, 1 H, J = 8.6 Hz), 7.09 - 7.04 (m, 3 H), 6.79 (dd, 1 H, J = 8.8 Hz), 5.11 (s, 2 H), 3.80 (s, 3 H), 2.33 (s, 3 H); IR (KBr) 3400, 2900, 1610  $\text{cm}^{-1}$ ; MS eI m/z 343 (M $^+$ ); CHN calcd for  $\text{C}_{23}\text{H}_{21}\text{NO}_2$  + 0.25  $\text{H}_2\text{O}$ .

**Example No. 5 5-methoxy-2-(4-methoxy-phenyl)-3-methyl-1H-indole**  
Mp = 139 - 142°C; <sup>1</sup>H NMR (DMSO) 10.85 (s, 1 H), 7.57 (d, 2 H, J = 8.8 Hz), 7.19 (d, 1 H, J = 8.6 Hz), 7.04 (d, 2 H, J = 6.8 Hz), 6.95 (d, 1 H, J = 2.2 Hz), 6.71 (dd, 1 H, J = 8.5 Hz, J = 2.4 Hz), 3.80 (s, 3 H), 3.76 (s, 3 H), 2.33 (s, 3 H);  
5 MS eI m/z 267 (M+); CHN calc for C<sub>17</sub>H<sub>17</sub>NO<sub>2</sub>.

**Example No. 6 5-Benzyl oxy-2-(4-ethoxy-phenyl)-3-methyl-1H-indole**  
Mp = 143-145°C; <sup>1</sup>H NMR (DMSO) 10.86 (s, 1H), 7.54 (d, 2 H, J = 8.5 Hz), 7.46 (d, 2 H, J = 7.3 Hz), 7.41-7.37 (m, 2 H), 7.32-7.30 (m, 1 H), 7.20 (d, 1 H, J = 8.6 Hz), 7.05 (d, 1 H), 7.03 (d, 2 H, J = 8.8 Hz), 6.79 (dd, 1 H, J = 8.6 Hz, J = 2.4 Hz), 5.10 (s, 2 H), 4.07 (q, 2 H, J = 6.8 Hz), 2.32 (s, 3 H), 1.34 (t, 3 H, J = 7.0 Hz); MS eI m/z 357 (M+).

**Example No. 8 5-Benzyl oxy-2-(4-fluoro-phenyl)-3-methyl-1H-indole**  
15 Mp = 132°C; <sup>1</sup>H NMR (DMSO) 11.0 (s, 1 H), 7.68-7.64 (m, 2 H), 7.49-7.47 (m, 2 H), 7.41-7.31 (m, 5 H), 7.23 (d, 1 H, J = 8.8 Hz), 7.10 (d, 1 H, J = 2.4 Hz), 6.82 (dd, 1 H, J = 8.8, 2.4 Hz), 5.11 (s, 2 H), 2.34 (s, 3 H); MS EI m/z 331; CHN calcd for C<sub>22</sub>H<sub>18</sub>FNO.

20 **Example No. 9 5-Benzyl oxy-2-(4-benzyl oxy-3-methoxy-phenyl)-3-methyl-1H-indole**  
Mp = 155 -158°C; <sup>1</sup>H NMR (DMSO) 10.88 (s, 1H), 7.50 - 7.45 (m, 4 H), 7.41 - 7.35 (m, 6H), 7.22 - 7.20 (m, 2 H), 7.14 (s, 2 H), 7.08 (d, 1 H, J = 2.2Hz), 6.78 (dd, 1H, J = 8.5 Hz, J = 2.4Hz), 5.13 (s, 2H), 5.11(s, 2H), 3.85 (s, 3H), 2.35 (s, 3H); MS eI m/z 449 (M+).

25 **Example No. 10 2-Benzof[1,3]dioxol-5-yl-5-benzyl oxy-3-methyl-1H-indole**  
Mp = 142-145°C; <sup>1</sup>H NMR (DMSO) 10.86 (s, 1H), 7.48 (d, 2 H, J = 7.0 Hz), 7.40 - 7.30 (m, 3 H), 7.20 (m, 2 H), 7.10 - 7.05 (m, 3 H), 6.78 (dd, 1 H, J = 8.8 Hz, J = 2.4 Hz), 6.06 (s, 2 H), 5.10 (s, 2 H), 2.31 (s, 3 H); MS eI m/z 357 (M+); CHN calc for C<sub>22</sub>H<sub>19</sub>NO<sub>3</sub>.

- 25 -

Example No. 11 5-Benzyl oxy-2-(4-isopropoxy-phenyl)-3-methyl-1H-indole

5 Mp = 136 - 138°C; <sup>1</sup>H NMR (DMSO) 10.86 (s, 1 H), 7.55 - 7.51 (m, 2 H), 7.50 - 7.47 (d, 2 H, J = 7.3 Hz), 7.40 - 7.34 (m, 2 H), 7.39 - 7.28 (m, 1 H), 7.20 (d, 1 H, J = 8.7 Hz), 7.06 (d, 1 H, J = 2.2 Hz), 7.02 (d, 2 H, J = 8.8 Hz), 6.77 (dd, 1 H, J = 2.4 Hz, 8.8 Hz), 5.10 (s, 2 H), 4.68 - 4.62 (m, 1 H), 2.32 (s, 3 H), 1.28 (d, 6 H, J = 6.0 Hz); MS eI m/z 371 (M+).

10 Example No. 12 5-Benzyl oxy-2-(4-cyclopropyl oxy-phenyl)-3-methyl-1H-indole

15 Mp = 161 - 167°C; <sup>1</sup>H NMR (DMSO) 10.85 (s, 1 H), 7.53 (d, 2 H, J = 8.8 Hz), 7.47 (d, 2 H, J = 8.4 Hz), 7.40 - 7.36 (m, 2 H), 7.33 - 7.28 (m, 1 H), 7.20 (d, 1 H, J = 8.6 Hz), 7.07 (d, 1 H, J = 2.4 Hz), 7.01 (d, 2 H, J = 8.8 Hz), 6.78 (dd, 1 H, J = 8.6 Hz, 2.2 Hz), 5.10 (s, 2 H), 4.88 - 4.84 (m, 1 H), 2.32 (s, 3 H), 1.99 - 1.88 (m, 2 H), 1.78 - 1.69 (m, 4 H), 1.64 - 1.52 (m, 2 H); IR (KBr) 3400, 2920, 1600 cm<sup>-1</sup>; MS eI m/z 397 (M+); CHN calcd for C<sub>27</sub>H<sub>27</sub>NO<sub>2</sub> + 0.25 H<sub>2</sub>O.

20 Example No. 13 5-Benzyl oxy-2-(4-trifluoromethyl-phenyl)-3-methyl-1H-indole

25 <sup>1</sup>H NMR (DMSO) 11.0 (br s, 1 H), 7.87 - 7.82 (m, 4 H), 7.48 (d, 2 H, J = 8.8 Hz), 7.44 - 7.35 (m, 2 H), 7.34 - 7.26 (m, 2 H), 7.15 (d, 1 H, J = 2.2 Hz), 6.87 (dd, 1 H, J = 8.6 Hz, 2.4 Hz), 5.12 (s, 2 H), 2.41 (s, 3 H); CHN calcd for C<sub>23</sub>H<sub>18</sub>F<sub>3</sub>NO.

30 Example No. 14 5-Benzyl oxy-2-(4-methyl-phenyl)-3-methyl-1H-indole

Mp = 144 - 146°C; <sup>1</sup>H NMR (DMSO) 10.91 (s, 1 H), 7.56 - 7.20 (m, 10 H), 7.08 (d, 1 H, J = 2.4 Hz), 6.80 (dd, 1 H, J = 2.4 Hz, 8.6 Hz), 5.11 (s, 2 H), 2.34 (s, 3 H), 2.34 (s, 3 H); MS eI m/z 327(M+).

35 Example No. 15 5-Benzyl oxy-2-(4-chloro-phenyl)-3-methyl-1H-indole

Mp = 134-136°C; <sup>1</sup>H NMR (DMSO) 11.04 (s, 1 H), 7.65 (d, 2 H, J = 8.3 Hz), 7.53

(d, 2 H, J = 8.5 Hz), 7.47 (d, 2 H, J = 6.8 Hz), 7.41 - 7.37 (m, 2 H), 7.31 - 7.28 (m, 1 H), 7.25 (d, 1 H, J = 8.5 Hz), 7.11 (d, 1 H, J = 2.4 Hz), 6.82 (dd, 1 H, J = 8.8 Hz, J = 2.4 Hz), 5.11 (s, 2 H), 2.35 (s, 3 H); IR (KBr) 3380, 1210  $\text{cm}^{-1}$ , MS eI m/z 347 (M+); CHN calc for  $\text{C}_{22}\text{H}_{18}\text{ClNO}_2$ .

5

Example No. 16 5-Benzyl oxy-2-(2,4-dimethoxy-phenyl)-3-methyl-1H-indole

Oil;  $^1\text{H}$  NMR (DMSO) 10.58 (s, 1 H), 7.50 - 7.18 (m, 7 H), 7.04 (d, 1 H, J = 2.4 Hz), 6.76 (dd, 1 H, J = 2.3 Hz, 8.6 Hz), 6.69 - 6.62 (m, 2 H), 5.11 (s, 2 H), 3.82 (s, 3 H), 3.78 (s, 3 H), 2.12 (s, 3 H).

10 Example No. 17 5-Benzyl oxy-2-(3-benzyl oxy-phenyl)-3-methyl-1H-indole

Mp = 83 - 86°C

15

Example No. 18 5-Benzyl oxy-2-(4-benzyl oxy-3-fluoro-phenyl)-3-methyl-1H-indole

Mp = 135 - 137°C;  $^1\text{H}$  NMR (DMSO) 10.94 (s, 1 H), 7.50 - 7.31 (m, 13 H), 7.22 (d, 1 H, J = 8.6 Hz), 7.10 (d, 1 H, J = 2.2 Hz), 6.81 (dd, 1 H, J = 8.6 Hz, 2.2 Hz), 5.23 (s, 2 H), 5.11 (s, 2 H), 2.34 (s, 3 H); MS eI m/z 437 (M+); CHN calcd for  $\text{C}_{29}\text{H}_{24}\text{FNO}_2$ .

20 Example No. 19 5-Benzyl oxy-2-(3-methoxy-phenyl)-3-methyl-1H-indole

Mp = 107 - 109°C;  $^1\text{H}$  NMR (DMSO) 11.00 (s, 1 H), 7.51 - 7.48 (m, 2 H), 7.43 - 7.20 (m, 7 H), 7.13 - 7.12 (d, 1 H, J = 2.1 Hz), 6.93 - 6.90 (dd, 1 H, J = 2.3 Hz, J = 5.7 Hz), 6.86 - 6.82 (dd, 1 H, J = 2.3 Hz, J = 6.3 Hz), 5.12 (s, 2 H), 3.83 (s, 3 H), 2.38 (s, 3 H); IR (KBr) 3400, 2900, 1600  $\text{cm}^{-1}$ ; MS eI m/z 343 (M+); CHN calcd for  $\text{C}_{21}\text{H}_{21}\text{NO}_2$ .

30

Example No. 20 5-Benzyl oxy-3-methyl-2-(4-trifluoromethoxy-phenyl)-1H-indole

Mp = 127 - 128°C;  $^1\text{H}$  NMR (DMSO) 11.07 (s, 1 H), 7.77 - 7.74

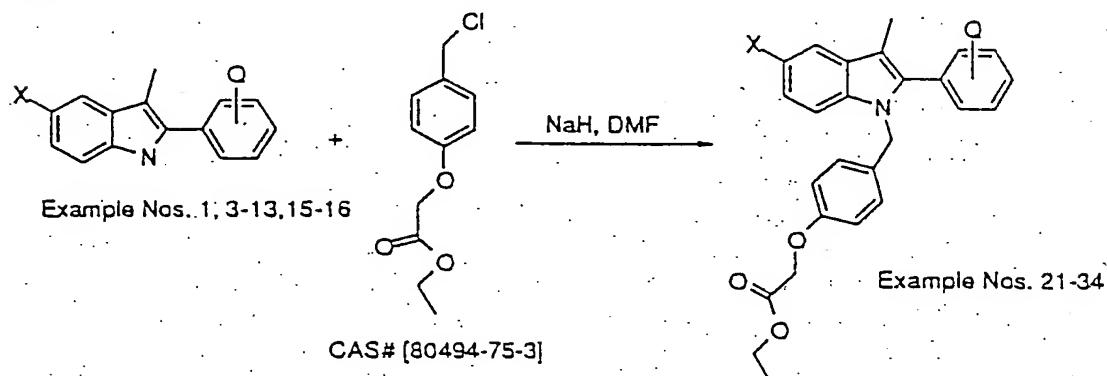
- 27 -

(dd, 2 H,  $J$  = 1.8 Hz,  $J$  = 5.0 Hz), 7.50 - 7.48 (d, 4 H,  $J$  = 8.3 Hz), 7.42 - 7.25 (m, 4 H), 7.14 - 7.13 (d, 1 H,  $J$  = 2.2 Hz), 6.87 - 6.83 (dd, 1 H,  $J$  = 2.3 Hz,  $J$  = 6.3 Hz), 5.13 (s, 2 H), 2.37 (s, 3 H); IR (KBr) 3360, 1600  $\text{cm}^{-1}$ ; MS eI m/z 396 (M $^+$ ); CHN calcd for  $\text{C}_{23}\text{H}_{18}\text{F}_3\text{NO}_2$ .

5

3-Methylindole acetic acid ethyl esters

Scheme 9



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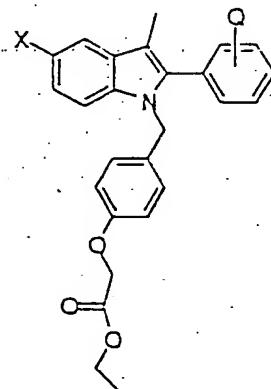


Table 2

Example No.	X	Q
No. 21	H	H
No. 22	OBn	H
No. 23	OBn	4'-OMe
No. 24	OMe	4'-OMe
No. 25	OBn	4'-OEt
No. 26	OBn	4'-OBn
No. 27	OBn	4'-F
No. 28	OBn	3'-OMe,4'-OBn
No. 29	OBn	4'-O-iPr
No. 30	OBn	3',4'-OCH <sub>2</sub> O-
No. 31	OBn	4'-OCp
No. 32	OBn	4'-CF <sub>3</sub>
No. 33	OBn	4'-Cl
No. 34	OBn	2'-OMe,4'-OMe

Experimental Procedure For 3-Methylindole Acetic Acid Ethyl Esters

5

Synthesis Method 3

Illustrated For Example No. 26

{4-[5-Benzylxy-2-(4-benzylxy-phenyl)-3-methyl-indol-1-yl]methoxy-phenoxyl}-acetic acid ethyl ester

10

A solution of 5-Benzylxy-2-(4-benzylxy-phenyl)-3-methyl-1H-indole (indole example No. 7) (32g, 77 mmol) in DMF (0.15 L) was cooled to 0°C and treated with sodium hydride (2.2 g, 89 mmol). The reaction was stirred for 20 minutes and then the benzyl chloride CAS No. [80494-75-3] (29g, 127 mmol) was added and the reaction stirred for 18 hours at room temperature. The reaction mixture was poured into water and extracted with ethyl acetate. The ethyl acetate was washed with brine and dried over magnesium sulfate. The ethyl acetate was concentrated and triturated with ether to

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obtain 21 g of a white solid. The filtrate was concentrated and triturated with ether to give an additional 7 g of white solid for a total yield of 28 g: Mp = 129-131°C; <sup>1</sup>H NMR (DMSO) 7.47 (d, 4 H, J = 7.2 Hz), 7.39 (q, 4 H, J = 7.9 Hz), 7.36-7.32 (m, 1 H), 7.29 (d, 2 H, J = 8.8 Hz), 7.19 (d, 1 H, J = 9.0 Hz), 7.13-7.09 (m, 4 H), 5 6.80 (dd, 1 H, J = 8.8, 2.4 Hz), 6.73 (s, 4 H), 5.16 (s, 2 H), 5.13 (s, 2 H), 5.11 (s, 2 H), 4.66 (s, 2 H), 4.11 (q, 2 H, J = 7.2 Hz), 2.15 (s, 3 H), 1.16 (t, 3 H, J = 7.2 Hz); MS eI m/z 612.

Physical Data For Indole Ethyl Esters

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The following indole alkylation products were prepared according to scheme 9 using method 3 with the appropriately substituted 3-methyl indole selected from (No. 1- No. 16) as the starting material.

15

Example No. 21 {4-[2-Phenyl-3-methyl-indol-1-ylmethyll-phenoxy}-acetic acid ethyl ester

Oil; <sup>1</sup>H NMR (DMSO) 7.57 - 7.30 (m, 7 H), 7.13 - 7.02 (m, 2 H), 6.77 - 6.70 (m, 4 H), 5.22 (s, 2 H), 4.65 (s, 2 H), 4.09 (q, 2 H, J = 7.2 Hz), 2.20 (s, 3 H), 1.15 (t, 3 H, J = 7.0 Hz); MS eI m/z 399 (M+).

20

Example No. 22 {4-[5-Benzyl-2-phenyl-3-methyl-indol-1-ylmethyll-phenoxy}-acetic acid ethyl ester

Oil; <sup>1</sup>H NMR (DMSO) 7.50 - 7.40 (m, 10 H), 7.22 (d, 1 H, J = 8.4 Hz), 7.14 (d, 1 H, J = 2.5 Hz), 6.83 (d, 1 H, J = 2.5 Hz), 6.72 (s, 4 H), 5.18 (s, 2 H), 5.11 (s, 2 H), 4.65 (s, 2 H), 4.10 (q, 2 H, J = 7.2 Hz), 2.16 (s, 3 H), 1.14 (t, 3 H, J = 7.0 Hz); MS eI m/z 505 (M+).

Example No. 23 {4-[5-Benzyl-2-(4-methoxy-phenyl)-3-methyl-indol-1-ylmethyll-phenoxy}-acetic acid ethyl ester

30 Mp = 90 - 96°C; <sup>1</sup>H NMR (DMSO) 7.47 (d, 2 H, J = 6.8 Hz), 7.41 - 7.37 (m, 2 H), 7.33 - 7.27 (m, 3 H), 7.19 (d, 1 H, J = 8.8 Hz), 7.12 (d, 1 H, J = 2.4 Hz), 7.03 (d, 2 H, J = 8.8 Hz), 6.80 (dd, 1 H, J = 8.8 Hz, 2.4 Hz), 6.74 (s, 4 H), 5.16

(s, 2 H), 5.11 (s, 2 H), 4.65 (s, 2 H), 4.11 (q, 2 H,  $J = 7.0$  Hz), 3.79 (s, 3 H), 2.15 (s, 3 H), 1.16 (t, 3 H,  $J = 7.0$  Hz); IR (KBr) 2990, 2900, 1760, 1610  $\text{cm}^{-1}$ ; MS FAB m/z 536 (M+H+).

5 Example No. 24 {4-[5-Methoxy-2-(4-methoxy-phenyl)-3-methyl-indol-1-ylmethyl-phenoxyl]-acetic acid ethyl ester}

Mp = 109-113°C;  $^1\text{H}$  NMR (DMSO) 7.27 (d, 2 H,  $J = 8.8$  Hz), 7.17 (d, 1 H,  $J = 8.8$  Hz), 7.03 (d, 2 H,  $J = 8.6$  Hz), 6.99 (d, 1 H,  $J = 2.5$  Hz), 6.78 - 6.70 (m, 5 H), 5.15 (s, 2 H), 4.65 (s, 2 H), 4.11 (q, 2 H,  $J = 7.0$  Hz), 3.78 (s, 3 H), 3.76 (s, 3 H), 2.15 (s, 3 H), 1.15 (t, 3 H,  $J = 7.1$  Hz); MS eI m/z 459 (M+).

Example No. 25 {4-[5-Benzyl oxy-2-(4-ethoxy-phenyl)-3-methyl-indol-1-ylmethyl-phenoxyl]-acetic acid ethyl ester}

15 Mp = 113-115°C;  $^1\text{H}$  NMR (DMSO) 7.45 (d, 2 H,  $J = 7.3$  Hz), 7.40 - 7.25 (m, 5 H), 7.17 (d, 1 H,  $J = 8.8$  Hz), 7.11 (d, 1 H,  $J = 2.2$  Hz), 7.01 (d, 2 H,  $J = 6.8$  Hz), 6.78 (dd, 1 H,  $J = 8.8$  Hz,  $J = 2.4$  Hz), 6.73 (s, 4 H), 5.15 (s, 2 H), 5.10 (s, 2 H), 4.65 (s, 2 H), 4.15 - 4.01 (m, 4 H), 2.14 (s, 3 H), 1.33 (t, 3 H,  $J = 5.7$  Hz), 1.16 (t, 3 H,  $J = 7.1$  Hz); MS eI m/z 549 (M+).

20

Example No. 27 {4-[5-Benzyl oxy-2-(4-fluoro-phenyl)-3-methyl-indol-1-ylmethyl-phenoxyl]-acetic acid ethyl ester}

$^1\text{H}$  NMR (DMSO) 7.50 - 7.15 (m, 16 H), 5.20 (s, 2 H), 5.12 (s, 2 H), 4.62 (s, 2 H), 4.13 (q, 2 H,  $J = 7.1$  Hz), 2.18 (s, 3 H), 1.20 (t, 3 H,  $J = 7.1$  Hz).

25

Example No. 28 {4-[5-Benzyl oxy-2-(3-methoxy-4-benzyl oxy)-3-methyl-indol-1-ylmethyl-phenoxyl]-acetic acid ethyl ester}

Foam;  $^1\text{H}$  NMR (DMSO) 7.50 - 7.30 (m, 10 H), 7.22 (d, 2 H,  $J = 9.1$  Hz), 7.13 (d, 2 H,  $J = 8.6$  Hz), 6.85 - 6.70 (m, 6 H), 5.17 (s, 2 H), 5.13 (s, 2 H), 5.11 (s, 2 H), 4.66 (s, 2 H), 4.14 (m, 2 H), 3.61 (s, 3 H), 2.17 (s, 3 H), 1.16 (t, 3 H,  $J = 7.0$  Hz).

30

Example No. 29 {4-[5-Benzylxy-2-(4-isopropoxy-phenyl)-3-methyl-indol-1-ylmethyll-phenoxy}-acetic acid ethyl ester

Oil;  $^1\text{H}$  NMR (DMSO) 7.46 (d, 2H,  $J = 7.7$  Hz), 7.42 - 7.28 (m, 3 H), 7.25 (d, 2 H,  $J = 8.7$  Hz), 7.17 (d, 1 H,  $J = 8.7$  Hz), 7.11 (d, 1 H,  $J = 2.4$  Hz), 6.99 (d, 2 H,  $J = 8.6$  Hz), 6.79 (dd, 1 H,  $J = 2.4$  Hz, 8.8 Hz), 6.73 (s, 4 H), 5.15 (s, 2 H), 5.10 (s, 2 H), 4.70 - 4.60 (m, 3 H), 4.10 (q, 2 H,  $J = 7.0$  Hz), 2.15 (s, 3 H), 1.27 (d, 6 H,  $J = 5.9$  Hz), 1.16 (t, 3 H,  $J = 7.1$  Hz); MS eI m/z 563 (M+).

Example No. 30 {4-[5-Benzylxy-2-(3,4-methylenedioxv-benzylxy)-3-methyl-indol-1-ylmethyll-phenoxy}-acetic acid ethyl ester

Oil;  $^1\text{H}$  NMR (DMSO) 7.45 (d, 2 H,  $J = 7.0$  Hz), 7.37 (m, 2 H), 7.32 (m, 1 H), 7.19 (d, 1 H,  $J = 8.8$  Hz), 7.11 (d, 1 H,  $J = 2.2$  Hz), 7.00 (d, 1 H,  $J = 7.9$  Hz), 6.90 (d, 1 H, 5.0 Hz), 6.82 - 6.75 (m, 6H), 6.07 (s, 2H), 5.16 (s, 2 H), 5.10 (s, 2H), 4.65 (s, 2 H), 4.10 (m, 2 H), 2.15 (s, 3 H), 1.15 (t, 3 H,  $J = 7.0$  Hz);  
MS eI m/z 549 (M+).

Example No. 31 {4-[5-Benzylxy-2-(4-cyclopentyloxy-phenyl)-3-methyl-indol-1-ylmethyll-phenoxy}-acetic acid ethyl ester

Mp = 96-98°C;  $^1\text{H}$  NMR (DMSO) 7.47 (d, 1 H,  $J = 7.2$  Hz), 7.40 - 7.36 (m, 2 H), 7.33 - 7.30 (m, 1 H), 7.26 (m, 2 H), 7.18 (d, 1 H,  $J = 8.8$  Hz), 7.11 (d, 1 H,  $J = 2.4$  Hz), 6.98 (d, 2 H,  $J = 8.8$  Hz), 6.79 (dd, 1 H,  $J = 8.8$  Hz, 2.4 Hz), 6.74 (s, 5 H), 5.15 (s, 2 H), 5.11 (s, 2 H), 4.86 - 4.80 (m, 1 H), 4.66 (s, 2 H), 4.13 (q, 2 H,  $J = 7.2$  Hz), 2.15 (s, 3 H), 1.98 - 1.85 (m, 2 H), 1.79 - 1.65 (m, 4 H), 1.62 - 1.55 (m, 2 H), 1.16 (t, 3 H,  $J = 7.0$  Hz); IR (KBr) 2950, 2910, 2890, 1760, 1610  $\text{cm}^{-1}$ ; MS eI m/z 589 (M+); CHN calcd for C:77.39 H:6.67 N: 2.38 Found: C:76.76 H:6.63 N:2.27.

Example No. 32 {4-[5-Benzylxy-3-methyl-2-(4-trifluoromethyl-phenyl)-indol-1-ylmethyll-phenoxy}-acetic acid ethyl ester

Mp = 221°C;  $^1\text{H}$  NMR (DMSO) 7.83 (d, 2 H,  $J = 8.1$  Hz), 7.60 (d, 2 H,  $J = 7.9$  Hz), 7.48 (d, 2 H,  $J = 8.4$  Hz), 7.40 - 7.36 (m, 4 H), 7.18 (d, 1 H,  $J = 2.4$  Hz), 6.86 (dd, 1 H,  $J = 8.8$  Hz, 2.4 Hz), 6.72 (s, 4 H), 5.21 (s, 2 H), 5.12 (s, 2 H), 4.65

(s, 2 H), 4.11 (q, 2 H,  $J = 7.2$  Hz), 2.20 (s, 3 H), 1.16 (t, 3 H,  $J = 7.0$  Hz); IR (KBr) 2920, 1730  $\text{cm}^{-1}$ ; MS eI m/z 573 (M $^+$ ); CHN calcd for  $\text{C}_{24}\text{H}_{30}\text{F}_3\text{NO}_4 + 0.25 \text{H}_2\text{O}$ .

Example No. 33 {4-[5-Benzylxoy-2-(4-chlorophenyl)-3-methyl-indol-1-ylmethyl-phenoxyl}-acetic acid ethyl ester

5 Mp = 99-101°C;  $^1\text{H}$  NMR (DMSO) 7.52 (d, 2 H,  $J = 8.6$  Hz), 7.46 (d, 2 H,  $J = 6.8$  Hz), 7.42 - 7.38 (m, 4 H), 7.36 (m, 1 H), 7.25 (d, 1 H,  $J = 9.0$  Hz), 7.14 (d, 1 H,  $J = 2.4$  Hz), 6.83 (dd, 1 H,  $J = 8.8$  Hz,  $J = 2.5$  Hz), 6.72 (s, 4 H), 5.18 (s, 2 H), 5.11 (s, 2 H), 4.65 (s, 2 H), 4.11 (q, 2 H,  $J = 7.2$  Hz), 2.16 (s, 3 H), 1.15 (t, 3 H,  $J = 7.2$  Hz); MS eI m/z 539 (M $^+$ ); CHN calc for  $\text{C}_{33}\text{H}_{30}\text{ClNO}_4$ .

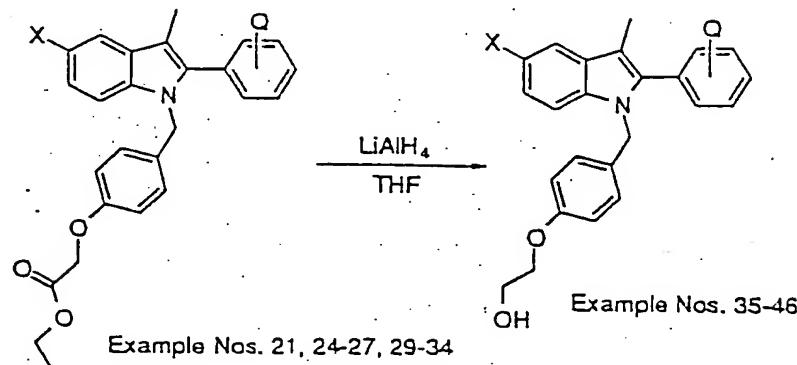
Example No. 34 {4-[5-Benzylxoy-2-(2,4-dimethoxy)-3-methyl-indol-1-ylmethyl-phenoxyl}-acetic acid ethyl ester

15 Oil;  $^1\text{H}$  NMR (DMSO) 7.30 - 6.45 (m, 15 H), 4.95 (s, 2 H), 4.75 - 4.65 (m, 2 H), 4.50 (s, 2 H), 3.97 (q, 2 H,  $J = 7.1$  Hz), 3.65 (s, 3 H), 3.51 (s, 3 H), 1.87 (3 H), 1.01 (t, 3 H,  $J = 7.1$  Hz).

20

### 3-Methylindole phenylethanols

Scheme 10



- 33 -

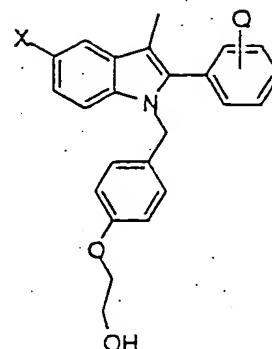


Table 3

Example No.	X	Q
No. 35	H	H
No. 36	OMe	4'-OMe
No. 37	OBn	4'-OEt
No. 38	OBn	4'-OBn
No. 39	OBn	4'-F
No. 40	OBn	3',4'-OCH <sub>2</sub> O-
No. 41	OBn	4'-O-iPr
No. 42	OBn	4'-OCp
No. 43	OBn	4'-CF <sub>3</sub>
No. 44	OBn	4'-CH <sub>3</sub>
No. 45	OBn	4'-Cl
No. 46	OBn	2'-OMe, 4'-OMe

5 Experimental Procedure For 3-Methylindole Phenethanols Synthesis  
Method 4

Illustrated For Example No. 38

10

2-(4-[5-Benzyl-2-(4-benzyloxy-phenyl)-3-methyl-indol-1-ylmethyl]-phenoxyl)-ethanol

A solution of No. 26 from previous step (5.5 g, 8.8 mmol) in THF (50 mL) was cooled to 0°C and a solution of LiAlH<sub>4</sub> (10mL, 1 M) in THF was added dropwise.

After 30 minutes at 0°C the reaction was carefully quenched with water, and partitioned between EtOAc and 1 N HCl. The EtOAc was dried with MgSO<sub>4</sub>, concentrated, and chromatographed on silica gel EtOAc/hexane (2:3) to yield 4.0 g of No. 38 as a white foam: <sup>1</sup>H NMR (DMSO) 7.48-7.46 (m, 4 H), 7.42-7.27 (m, 8 H), 7.20

5 (d, 1 H, J = 8.8 Hz), 7.12-7.10 (m, 3 H), 6.80 (dd, 1 H, J = 8.8, 2.4 Hz), 6.73 (s, 4 H), 5.15 (s, 2 H), 5.13 (s, 2 H), 5.11 (s, 2 H), 4.80 (t, 1 H, J = 5.5 Hz), 3.86 (t, 2 H, J = 4.8 Hz), 3.63 (q, 2 H, J = 5.3 Hz), 2.15 (s, 3 H).

#### Physical Data For Indole Phenethanols

10

Following compounds were made according to scheme 10 and method 4 using the appropriately substituted indole ethyl ester selected from No. 21- No. 34:

15 Example No. 35 2-[4-[2-phenyl-3-methyl-indol-1-ylmethyl]-phenoxy]-ethanol

Oil; <sup>1</sup>H NMR (DMSO) 7.57 - 7.32 (m, 7 H), 7.13 - 7.02 (m, 2 H), 6.74 (s, 4 H), 5.21 (s, 2 H), 4.80 (s, 1 H), 3.86 - 3.83 (m, 2 H), 3.62 (s, 2 H), 2.20 (s, 3 H); MS eI m/z 357 (M+).

20 Example No. 36 2-[4-[5-methoxy-2-(4-methoxy-phenyl)-3-methyl-indol-1-ylmethyl]-phenoxy]-ethanol

Oil; <sup>1</sup>H NMR (DMSO) 7.27 (d, 2 H, J = 8.8 Hz), 7.17 (d, 1 H, J = 8.8 Hz), 7.03 (d, 2 H, J = 8.6 Hz), 6.99 (d, 1 H, J = 2.5 Hz), 6.78 - 6.70 (m, 5 H), 5.14 (s, 2 H), 4.80 (brs, 1 H), 3.85 (t, 2 H, J = 5.0 Hz), 3.78 (s, 3 H), 3.76 (s, 3 H), 3.63 (t, 2 H, J = 5.0 Hz), 2.16 (s, 3 H); MS eI m/z 417 (M+).

Example No. 37 2-[4-[5-benzyl-2-(4-ethoxy-phenyl)-3-methyl-indol-1-ylmethyl]-phenoxy]-ethanol

Foam; <sup>1</sup>H NMR (DMSO) 7.45 (d, 2 H, J = 7.3 Hz), 7.40 - 7.25 (m, 5 H), 7.17 (d, 1 H, J = 8.8 Hz), 7.11 (d, 1 H, J = 2.2 Hz), 7.01 (d, 2 H, J = 6.8 Hz), 6.78 (dd, 1 H, J = 8.8 Hz, J = 2.4 Hz), 6.73 (s, 4 H), 5.15 (s, 2 H), 5.10 (s, 2 H), 4.80 (brs, 1 H), 4.06 (q, 2 H, J = 6.8 Hz), 3.85 (t, 2 H, J = 5.0 Hz), 3.63 (t, 2 H, J = 4.8 Hz), 2.14 (s, 3 H), 1.33 (t, 3 H, J = 6.9 Hz); MS eI m/z 507 (M+).

Example No. 39 2-[4-[5-benzyl oxy-2-(4-fluoro-phenyl)-3-methyl-indol-1-ylmethyl]-phenoxy]-ethanol

1H NMR (DMSO) 7.40 - 6.60 (m, 16 H), 5.10 (s, 1 H), 5.07 (s, 2 H), 5.02 (s, 2 H),  
5 3.76 (t, 2 H, J = 4.9 Hz), 3.53 (t, 2 H, J = 5.0 Hz), 2.06 (s, 3 H).

Example No. 40 2-[4-[5-benzyl oxy-2-(3,4-methylenedioxo-phenyl)-3-methyl-indol-1-ylmethyl]-phenoxy]-ethanol

Oil; 1H NMR (DMSO) 7.45 (d, 2 H, J = 7.0 Hz), 7.37 (m, 2 H), 7.32 (m, 1 H),  
10 7.19 (d, 1 H, J = 8.8 Hz), 7.11 (d, 1 H, J = 2.2 Hz), 7.00 (d, 1 H, J = 7.9 Hz), 6.90  
(d, 1 H, 5.0 Hz), 6.82 - 6.75 (m, 6H), 6.07 (s, 2 H), 5.16 (s, 2 H), 5.10  
(s, 2 H), 3.86 (t, 2 H, J = 5.0 Hz), 3.63 (t, 2 H, J = 5.0 Hz), 2.15 (s, 3 H); MS eI  
m/z 507 (M+).

15 Example No. 41 2-[4-[5-Benzyl oxy-2-(4-isopropoxy-phenyl)-3-methyl-indol-1-ylmethyl]-phenoxy]-ethanol

Foam; 1H NMR (DMSO) 7.46 (d, 2H, J = 7.7 Hz), 7.42 - 7.28 (m, 3 H), 7.25  
(d, 2 H, J = 8.7 Hz), 7.17 (d, 1 H, J = 8.7 Hz), 7.11 (d, 1 H, J = 2.4 Hz), 6.99  
(d, 2 H, J = 8.6 Hz), 6.79 (dd, 1 H, J = 2.4 Hz, 8.8 Hz), 6.73 (s, 4 H), 5.14  
20 (s, 2 H), 5.10 (s, 2 H), 4.80 (bs, 1 H), 4.70 - 4.60 (m, 1 H), 3.85  
(t, 2 H, J = 4.8 Hz), 3.63 (t, 2 H, J = 5.1 Hz), 2.13 (s, 3 H), 1.30  
(d, 6 H, J = 5.9 Hz); MS eI m/z 521 (M+).

25 Example No. 42 2-[4-[5-Benzyl oxy-2-(4-cyclopentyl oxy-phenyl)-3-methyl-indol-1-ylmethyl]-phenoxy]-ethanol

Mp = 129-131°C; 1H NMR (DMSO) 7.47 (d, 2 H, J = 7.2 Hz), 7.38  
(t, 2 H, J = 7.2 Hz), 7.33 - 7.28 (m, 1 H), 7.25 (d, 2 H, J = 8.8 Hz), 7.18  
(d, 1 H, J = 8.8 Hz), 7.11 (d, 1 H, J = 2.4 Hz), 6.98 (d, 2 H, J = 8.8 Hz), 6.79  
(dd, 1 H, J = 8.8 Hz, 2.4 Hz), 6.74 (s, 4H), 5.15 (s, 2 H), 5.11 (s, 2 H), 4.84 - 4.80  
30 (m, 1 H), 4.79 (t, 1 H, J = 5.7 Hz), 3.86 (t, 2 H, J = 4.8 Hz), 3.63  
(q, 2 H, J = 5.1 Hz), 2.15 (s, 3 H), 1.96 - 1.87 (m, 2 H), 1.77 - 1.65 (m, 4 H), 1.62  
- 1.53 (m, 2 H); IR (KBr) 3490 br, 2920, 1620 cm-1; MS eI m/Z 547 (M+).

Example No. 43 2-[4-[5-Benzyl oxy-2-(4-trifluoromethyl-phenyl)-3-methyl-indol-1-ylmethyll-phenoxy]-ethanol

Foam;  $^1\text{H}$  NMR (DMSO) 7.83 (d, 2 H,  $J = 8.1$  Hz), 7.59 (d, 2 H,  $J = 7.9$  Hz), 7.47 (d, 2 H,  $J = 8.3$  Hz), 7.42 - 7.36 (m, 2 H), 7.35 - 7.29 (m, 2 H), 7.18 5 (d, 1 H,  $J = 2.4$  Hz), 6.87 (dd, 1 H,  $J = 8.1$  Hz, 2.4 Hz), 6.77 - 6.68 (m, 4 H), 5.21 (s, 2 H), 5.12 (s, 2 H), 4.81 (br s, 1 H), 3.85 (t, 2 H,  $J = 5.1$  Hz), 3.63 (t, 2 H,  $J = 5.1$  Hz), 2.19 (s, 3 H); MS eI m/z 531.

Example No. 44 2-[4-[5-Benzyl oxy-2-(4-methyl-phenyl)-3-methyl-indol-1-ylmethyll-phenoxy]-ethanol

Oil;  $^1\text{H}$  NMR (DMSO) 7.46 (d, 2 H,  $J = 7.2$  Hz), 7.45 - 7.18 (m, 8 H), 7.12 (d, 1 H,  $J = 2.4$  Hz), 6.81 (dd, 1 H,  $J = 2.4$  Hz, 8.6 Hz), 6.73 (s, 4 H), 5.15 (s, 2 H), 5.10 (s, 2 H), 4.80 (bs, 1 H), 3.85 (t, 2 H,  $J = 4.8$  Hz), 3.63 (t, 2 H,  $J = 4.9$  Hz), 2.34 (s, 3 H), 2.15 (s, 3 H); MS eI m/z 477 (M $^+$ ).

Example No. 45 2-[4-[5-Benzyl oxy-2-(4-chloro-phenyl)-3-methyl-indol-1-ylmethyll-phenoxy]-ethanol

Mp = 110 - 113°C;  $^1\text{H}$  NMR (DMSO) 7.52 (d, 2 H,  $J = 8.6$  Hz), 7.46 (d, 2 H,  $J = 6.8$  Hz), 7.38 (m, 4 H), 7.42 - 7.37 (m, 1 H), 7.25 (d, 1 H,  $J = 9.0$  Hz), 7.14 (d, 1 H,  $J = 2.4$  Hz), 6.83 (dd, 1 H,  $J = 8.8$  Hz,  $J = 2.5$  Hz), 6.76 - 6.70 (m, 4 H), 5.17 (s, 2 H), 5.11 (s, 2 H), 3.85 (t, 2 H,  $J = 5.2$  Hz), 3.63 (t, 2 H,  $J = 5.0$  Hz), 2.16 (s, 3 H); MS eI m/z 497 (M $^+$ ).

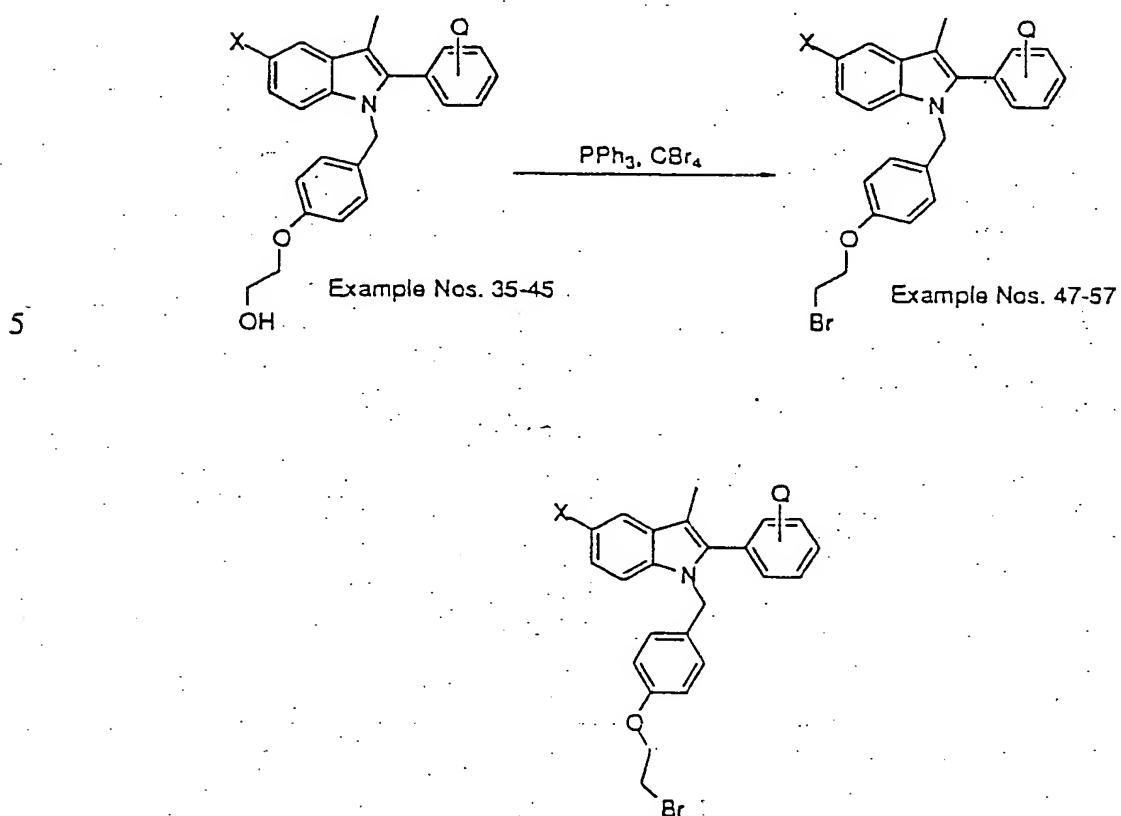
Example No. 46 2-[4-[5-Benzyl oxy-2-(2,4-dimethoxy-phenyl)-3-methyl-indol-1-ylmethyll-phenoxy]-ethanol

Oil;  $^1\text{H}$  NMR (DMSO) 7.46 (d, 2 H,  $J = 7.5$  Hz), 7.39 - 7.35 (m, 2 H), 7.31 - 7.28 (m, 1 H), 7.16 - 7.06 (m, 3 H), 6.82 - 6.72 (m, 5 H), 6.68 (d, 1 H,  $J = 2.2$  Hz), 6.61 (dd, 1 H,  $J = 2.4$  Hz, 8.3 Hz), 5.0 (s, 1 H), 4.88 (s, 2 H), 4.85 (d, 1 H,  $J = 6.3$  Hz), 4.69 (d, 1 H,  $J = 6.3$  Hz), 3.63 (t, 2 H,  $J = 6.9$  Hz), 3.58 (s, 3 H), 3.46 (s, 3 H), 30 3.40 (t, 2 H,  $J = 6.9$  Hz), 1.80 (s, 3 H).

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Data for 3-methylindole phenylethyl bromides

Scheme 11



Example No.	X	Q
No. 47	H	H
No. 48	OMe	4'-OMe
No. 49	OBn	4'-OEt
No. 50	OBn	4'-OBn

Table 4 (Cont'd)

Example No.	X	Q
No. 51	OBn	4'-F
No. 52	OBn	3',4'-OCH <sub>2</sub> O-
No. 52a	OBn	3'-OMe, 4'-OBn
No. 53	OBn	4'-O-iPr
No. 54	OBn	4'-OCp
No. 55	OBn	4'-CF <sub>3</sub>
No. 56	OBn	4'-CH <sub>3</sub>
No. 57	OBn	4'-Cl

5      Experimental Procedure For 3-Methylindole Phenethyl bromide  
Synthesis  
Method 5  
Illustrated For Example No. 50

10

Example No. 50

5-Benzyl oxy-2-(4-Benzyl oxy-phenyl)-1-[4-(2-bromo-ethoxy)-benzyl]-3-methyl-1H-indole

15      To a solution of example No. 38 (3.3 g, 5.8 mmol) in THF (50 mL) was added CBr<sub>4</sub> (2.9 g, 8.7 mmol) and PPH<sub>3</sub> (2.3 g, 8.7 mmol). The reaction was stirred at  $\pi$  for 3 h and then concentrated and chromatographed on silica gel using a gradient elution from EtOAc/hexane (1:4) to EtOAc to give 3.2 g of a white solid: Mp = 131-134°C; <sup>1</sup>H NMR (DMSO) 7.64-7.30 (m, 10 H), 7.29 (d, 2 H, J = 8.8 Hz), 7.20 (d, 1 H, J = 8.8 Hz), 7.12-7.09 (m, 3 H), 6.80 (dd, 1 H, J = 8.8, 2.4 Hz), 6.77-6.73 (m, 4 H), 5.16 (s, 2 H), 5.13 (s, 2 H), 5.11 (s, 2 H), 4.20 (t, 2 H, J = 5.3 Hz), 3.73 (t, 2 H, J = 5.5 Hz), 2.15 (s, 3 H); MS FAB 631/633 (M+H<sup>+</sup>, Br present).

### Physical Data for Indole Phenethyl Bromides

The following compounds were made according to scheme 11 as described in Method 5 using the appropriately substituted indole selected from No. 35- No. 45.

5

#### Example No. 47 1-[4-(2-bromo-ethoxy)-benzyl]-2-phenyl-3-methyl-1H-indole

Oil;  $^1\text{H}$  NMR (DMSO) 7.57 - 7.32 (m, 7 H), 7.13 - 7.02 (m, 2 H), 6.74 (s, 4 H), 10 5.21 (s, 2 H), 4.19 (t, 2 H,  $J$  = 5.2 Hz), 3.71 (t, 2 H,  $J$  = 5.5 Hz), 2.20 (s, 3 H); MS eI m/z 419 (M+).

#### Example No. 48 5-Methoxy-2-(4-methoxy-phenyl)-1-[4-(2-bromo-ethoxy)-benzyl]-3-methyl-1H-indole

Oil;  $^1\text{H}$  NMR (DMSO) 7.27 (d, 2 H,  $J$  = 8.8 Hz), 7.17 (d, 1 H,  $J$  = 8.8 Hz), 7.03 (d, 2 H,  $J$  = 8.6 Hz), 6.99 (d, 1 H,  $J$  = 2.5 Hz), 6.80 - 6.69 (m, 5 H), 5.14 (s, 2 H), 4.19 (t, 2 H,  $J$  = 5.4 Hz), 3.78 (s, 3 H), 3.76 (s, 3 H), 3.72 (t, 2 H,  $J$  = 5.5 Hz), 2.16 (s, 3 H); MS eI m/z 479 (M+).

#### Example No. 49 5-Benzyl oxy-2-(4-ethoxy-phenyl)-1-[4-(2-bromo-ethoxy)-benzyl]-3-methyl-1H-indole

Mp = 118-120°C;  $^1\text{H}$  NMR (DMSO) 7.45 (d, 2 H,  $J$  = 7.3 Hz), 7.41 - 7.26 (m, 5 H), 7.17 (d, 1 H,  $J$  = 8.8 Hz), 7.11 (d, 1 H,  $J$  = 2.2 Hz), 7.01 (d, 2 H,  $J$  = 6.8 Hz), 6.78 (dd, 1 H,  $J$  = 8.8 Hz,  $J$  = 2.4 Hz), 6.78 - 6.74 (m, 4 H), 25 5.15 (s, 2 H), 5.10 (s, 2 H), 4.22 - 4.18 (m, 2 H), 4.04 (q, 2 H,  $J$  = 6.8 Hz), 3.72 (t, 2 H,  $J$  = 5.5 Hz), 2.14 (s, 3 H), 1.33 (t, 3 H,  $J$  = 7.0 Hz); MS eI m/z 569 (M+).

#### Example No. 51 5-Benzyl oxy-1-[4-(2-bromo-ethoxy)-benzyl]-2-(4-fluoro-phenyl)-3-methyl-1H-indole

30 Mp = 114-116°C;  $^1\text{H}$  NMR (DMSO) 7.47 (m, 2 H), 7.45 - 7.20 (m, 8 H), 7.14 (d, 1 H,  $J$  = 2.4 Hz), 6.83 (dd, 1 H,  $J$  = 2.7 Hz, 9.0 Hz), 6.80 - 6.70 (m, 4 H), 5.16 (s, 2 H), 5.11 (s, 2 H), 4.19 (t, 2 H,  $J$  = 5.27 Hz), 3.72 (t, 2 H,  $J$  = 6.4 Hz), 2.15 (s, 3 H); MS eI m/z 543 (M+); CHN calc for  $\text{C}_{31}\text{H}_{27}\text{BrFNO}_2$ .

Example No. 52 2-Benzof[1,3]dioxyl-5-vl-5-benzylloxv-1-[4-(2-bromo-ethoxy)-benzyl]-3-methyl-1H-indole

5 Mp = 133-136°C; <sup>1</sup>H NMR (DMSO) 7.45 (d, 2 H, J = 7.0 Hz), 7.41-7.38 (m, 2 H), 7.35-7.30 (m, 1 H), 7.19 (d, 1 H, J = 8.8 Hz), 7.11 (d, 1 H, J = 2.2 Hz), 7.00 (d, 1 H, J = 7.9 Hz), 6.90 (d, 1 H, 1.4 Hz), 6.82 - 6.78 (m, 2H), 6.77 (s, 4 H), 6.07 (s, 2 H), 5.16 (s, 2 H), 5.10 (s, 2 H), 4.20 (t, 2 H, J = 5.5Hz), 3.73 (t, 2H, J = 5.2Hz), 2.15 (s, 3H); MS eI m/z 569 (M+).

10 Example No. 52a 5-Benzylloxv-1-[4-(2-bromo-ethoxy)-benzyl]-2-(3-methoxy-4-benzylloxv-phenyl)-3-methyl-1H-indole

15 Foam; <sup>1</sup>H NMR (DMSO) 7.47 - 7.42 (m, 4 H), 7.40 - 7.30 (m, 6 H), 7.20 (d, 1 H, J = 8.8 Hz), 7.12 - 7.10 (m, 2 H), 6.86 - 6.84 (m, 2 H), 6.81 (dd, 1 H, J = 8.8 Hz, 2.4 Hz), 6.78 (s, 4 H), 5.17 (s, 2 H), 5.11 (s, 2 H), 5.10 (s, 2 H), 4.20 (t, 2 H, J = 5.0 Hz), 3.72 (t, 2 H, J = 5.4 Hz), 3.63 (s, 3 H), 2.17 (s, 3 H); MS FAB m/z 662 (M+H+).

Example No. 53 5-Benzylloxv-1-[4-(2-bromo-ethoxy)-benzyl]-2-(4-isopropoxy-phenyl)-3-methyl-1H-indole

20 Mp = 125 - 128°C; <sup>1</sup>H NMR (DMSO) 7.46 (d, 2H, J = 7.7 Hz), 7.42 - 7.28 (m, 3 H), 7.25 (d, 2 H, J = 8.7 Hz), 7.17 (d, 1 H, J = 8.7 Hz), 7.11 (d, 1 H, J = 2.4 Hz), 6.99 (d, 2 H, J = 8.6 Hz), 6.79 (dd, 1 H, J = 2.4 Hz, 8.8 Hz), 6.73 (s, 4 H), 5.14 (s, 2 H), 5.10 (s, 2 H), 4.70 - 4.60 (m, 1 H), 4.19 (t, 2 H, J = 5.3 Hz), 3.72 (t, 2 H, J = 4.4 Hz), 2.13 (s, 3 H), 1.30 (d, 6 H, J = 5.9 Hz); MS eI m/z 583 (M+).

25

Example No. 54 5-Benzyloxy-1-[4-(2-bromo-ethoxy)-benzyl]-2-(4-cyclopentyloxv-phenyl)-3-methyl-1H-indole

30 Mp = 110 - 112°C; 7.47 (d, 2 H, J = 7.0 Hz), 7.38 (t, 2 H, J = 7.0 Hz), 7.35 - 7.28 (m, 1 H), 7.25 (d, 2 H, J = 8.8 Hz), 7.18 (d, 1 H, J = 8.8 Hz), 7.11 (d, 1 H, J = 2.4 Hz), 6.98 (d, 2 H, J = 8.6 Hz), 6.79 (dd, 1 H, J = 8.6 Hz, 2.4 Hz), 6.78 - 6.74 (m, 4 H), 5.16 (s, 2 H), 5.11 (s, 2 H), 4.86 - 4.83 (m, 1 H), 4.20

(t, 2 H, J = 5.3 Hz), 3.73 (t, 2 H, J = 5.5 Hz), 2.15 (s, 3 H), 2.00 - 1.87 (m, 2 H), 1.79 - 1.65 (m, 4 H), 1.63 - 1.56 (m, 2 H); IR (KBr) 2950, 2910, 1610 cm<sup>-1</sup>; MS eI m/z 609, 611 (M+, Br present).

5 Example No. 55 5-Benzyl oxy-1-[4-(2-bromo-ethoxy)-benzyl]-3-methyl-2-(4-trifluoromethyl-phenyl)-1H-indole

Mp = 106 -109°C; <sup>1</sup>H NMR (DMSO) 7.83 (d, 2 H, J = 8.1 Hz), 7.60 (d, 2 H, J = 7.9 Hz), 7.35 - 7.29 (m, 2 H), 7.48 (d, 2 H, J = 8.6 Hz), 7.39 (t, 2 H, J = 7.0 Hz), 7.18 (d, 1 H, J = 2.2 Hz), 6.87 (dd, 1 H, J = 9.0 Hz, 2.6 Hz), 10 6.77 - 6.71 (m, 4 H), 5.22 (s, 2 H), 5.12 (s, 2 H), 4.20 (t, 2 H, J = 5.3 Hz), 3.72 (t, 2 H, J = 5.3 Hz), 2.20 (s, 3 H); IR (KBr) 2910, 2850, 1620 cm<sup>-1</sup>; MS eI m/z 595, 593 (M+)

15 Example No. 56 5-Benzyl oxy-1-[4-(2-bromo-ethoxy)-benzyl]-3-methyl-2-(4-methyl-phenyl)-1H-indole

Mp = 82 - 95°C; <sup>1</sup>H NMR (DMSO) 7.46 (d, 2 H, J = 7.2 Hz), 7.45 - 7.18 (m, 8 H), 7.12 (d, 1 H, J = 2.4 Hz), 6.81 (dd, 1 H, J = 2.4 Hz, 8.6 Hz), 6.73 (s, 4 H), 5.15 (s, 2 H), 5.10 (s, 2 H), 4.19 (t, 2 H, J = 5.3 Hz), 3.72 (t, 2 H, J = 4.4 Hz), 2.34 (s, 3 H), 2.15 (s, 3 H); MS eI m/z 539 (M+).

20

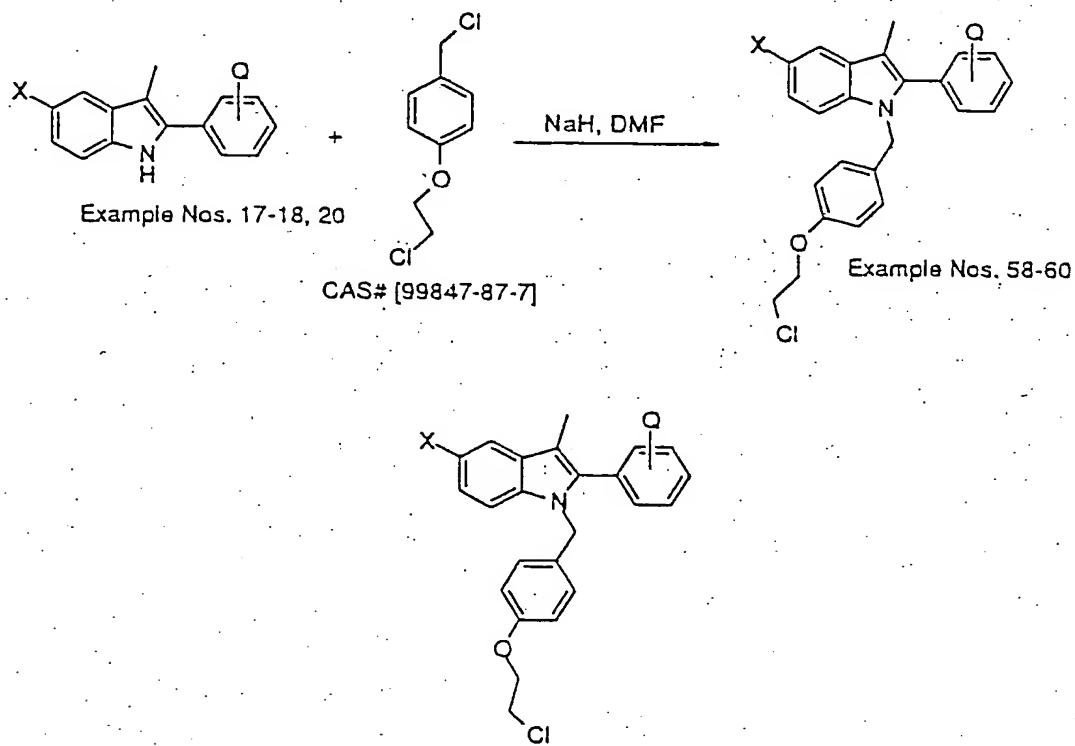
Example No. 57 5-Benzyl oxy-1-[4-(2-bromo-ethoxy)-benzyl]-3-methyl-2-(4-chloro-phenyl)-1H-indole

<sup>1</sup>H NMR (DMSO) 7.52 (d, 2 H, J = 8.6 Hz), 7.46 (d, 2 H, J = 6.8 Hz), 7.38 (m, 4 H), 7.36 (m, 1 H), 7.25 (d, 1 H, J = 9.0 Hz), 7.14 (d, 1 H, J = 2.4 Hz), 6.83 (dd, 1 H, J = 8.8 Hz, J = 2.5 Hz), 6.72 (m, 4 H), 5.17 (s, 2 H), 5.11 (s, 2 H), 4.19 (t, 2 H, J = 5.5 Hz), 3.72 (t, 2 H, J = 5.5 Hz), 2.16 (s, 3 H); MS eI m/z 559 (M+).

Data for some 3-methylindole phenylethyl chlorides used as intermediates

Scheme 12

5



10

Table 5

Example No.	X	Q
No. 58	OBn	3'-OBn
No. 59	OBn	3'-F, 4'-OBn
No. 60	OBn	4'-OCF <sub>3</sub>

Experimental Procedure For 3-Methylindole Phenethylchloride  
Synthesis  
Method 5a

5

Illustrated For Example No. 58

5-Benzylxy-2-(3-benzylxy-phenyl)-1-[4-(2-chloro-ethoxy)-benzyl-3-methyl-1H-indole

To a solution of 9.7 g (0.0231 mol) of 5-benzylxy-3-methyl-2-(3-benzylxy-phenyl)-1H-indole (indole example No. 17) in 80 mL of dry DMF was added 0.85 g 10 of sodium hydride (60% in mineral oil). After allowing this mixture to stir for 30 minutes (until no more bubbling was indicated), 4.8 g of 1-chloromethyl-4-(2-chloroethoxy)-benzene CAS No. [99847-87-7] was added. The reaction mixture was allowed to react at room temperature overnight. 200 mL of ethyl acetate were added to the reaction mixture and then washed with water (3 x 100 mL). The organic solution 15 was collected, washed with saturated brine, removed, dried over magnesium sulfate, filtered and evaporated to dryness in a rotary evaporator. The product was recrystallized in ethyl acetate.

Mp = 125 - 127°C; <sup>1</sup>H NMR (DMSO) 7.48 - 7.46 (d, 2 H, J = 6.8 Hz), 7.40 - 7.35 (m, 7 H), 7.33 - 7.28 (m, 2 H), 7.23 - 7.21 (d, 1 H, J = 8.8 Hz), 7.13 - 7.12 (d, 1 H, J = 2.2 Hz), 7.07 - 7.04 (m, 1 H), 6.94 - 6.92 (d, 2 H, J = 6.1 Hz), 6.83 - 6.80 (dd, 1 H, J = 2.5 Hz, J = 6.3 Hz), 6.78 - 6.72 (m, 4 H), 5.14 (s, 2 H), 5.11 (s, 2 H), 5.04 (s, 2 H), 4.13 - 4.10 (t, 2 H, J = 5.1 Hz), 3.86 - 3.84 (t, 2 H, J = 5.1 Hz), 2.14 (s, 3 H); IR 3420, 2900  $\text{cm}^{-1}$ ; MS eI m/z 587 (M+); CHN calcd for  $\text{C}_{18}\text{H}_{34}\text{ClNO}_3$ .

25

Physical Data for Indole Phenethyl Chlorides

The following compounds were made according to scheme 12 as described in Method 5a using the appropriately substituted indoles No. 18, No. 20.

30

**Example No. 59 5-Benzyl-2-(4-benzyl-3-fluoro-phenyl)-1-[4-(2-chloro-ethoxy)-benzyl-3-methyl-1H-indole**

5 Mp = 88-91°C;  $^1\text{H}$  NMR (DMSO) 7.49-7.43 (m, 4H), 7.43-7.28 (m, 7H), 7.26-7.21 (m, 2H), 7.13-7.09 (m, 2H), 6.88-6.72 (m, 5H), 5.21 (s, 2H), 5.18 (s, 2H), 5.11 (s, 2H), 4.13 (t, 2H,  $J = 5.2$  Hz), 3.87 (t, 2H,  $J = 5.2$  Hz), 2.16 (s, 3H); MS ei m/z 605 (M+); CHN calcd for  $\text{C}_{38}\text{H}_{33}\text{ClFNO}_3$ .

**Example No. 60 5-Benzyl-1-[4-(2-chloro-ethoxy)-benzyl-3-methyl-2-(4-trifluoromethoxy-phenyl)-1H-indole**

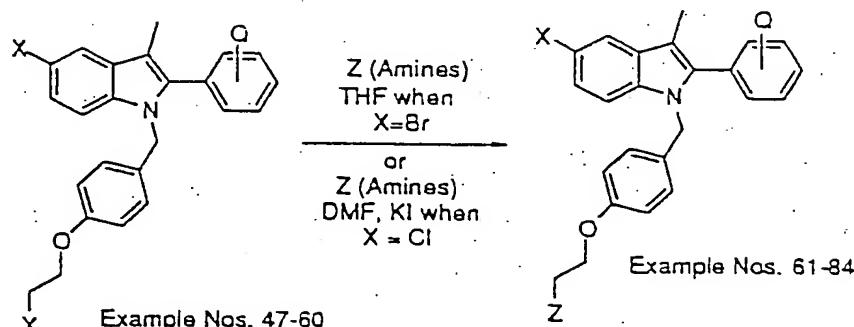
10 Mp = 108 - 110°C;  $^1\text{H}$  NMR (DMSO) 7.49 - 7.43 (m, 6 H), 7.40 - 7.25 (m, 4 H), 7.17 - 7.16 (d, 1 H,  $J = 2.9$  Hz), 6.88 - 6.84 (m, 1 H), 6.77 - 6.72 (m, 4 H), 5.20 (s, 2 H), 5.14 - 5.13 (d, 2 H,  $J = 2.3$  Hz), 4.16 - 4.11 (m, 2 H), 3.89 - 3.84 (m, 2 H), 2.19 - 2.17 (m, 3 H); IR 3400, 2900, 1600  $\text{cm}^{-1}$ ; MS ei m/z 566 (M+); CHN calcd for  $\text{C}_{32}\text{H}_{27}\text{ClF}_3\text{NO}_3 + 0.25 \text{H}_2\text{O}$ .

15

Aminoethoxy indoles

**Scheme 13**

20



- 45 -

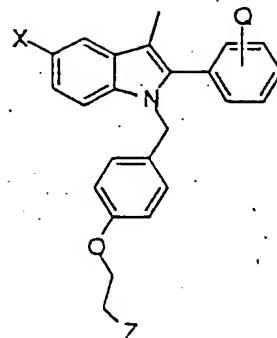


Table 6

5

Example No.	X	Q	Z
No. 61	OBn	4'-OEt	
No. 62	OBn	H	
No. 63	OBn	4'-OBn	
No. 64	OBn	4'-OBn	
No. 65	OBn	4'-OBn	

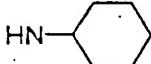
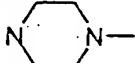
Table 6 (Cont'd)

Example No.	X	Q	Z
No. 66	OBn	4'-OBn	
No. 66a	OBn	4'-OBn	
No. 67	OBn	4'-OBn	
No. 68	OBn	4'-OBn	
No. 69	OBn	4'-OBn	
No. 70	OBn	4'-OBn	
No. 71	OBn	4'-OBn	
No. 71a	OBn	4'-OBn	
No. 72	OBn	4'-F	

Table 6 (Cont'd)

Example No.	X	Q	Z
No. 72a	OBn	4'-F	
No. 72b	OBn	4'-Cl	
No. 73	OBn	3',4'-OCH <sub>2</sub> O-	
No. 74	OBn	4'-O-iPr	
No. 75	OBn	4'-CH <sub>3</sub>	
No. 76	OBn	3'-OBn	
No. 77	OBn	3'-OBn	
No. 78	OBn	4'-OBn,3'-F	
No. 79	OBn	4'-OBn,3'-F	

Table 6 (Cont'd)

Example No.	X	Q	Z
No. 80	OBn	3'-OMe	
No. 81	OBn	4'-OCF <sub>3</sub>	
No. 82	OBn	4'-OBn	
No. 83	OBn	4'-OBn	
No. 84	OBn	3'-OMe	

Experimental Procedure For 3-Methyl aminoethoxyindole Synthesis

5.

Method 6

Illustrated For Example No. 63

Substitution of the Bromide

10 5-Benzyl oxy-2-(4-benzyl oxy-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole

A solution of example No. 50 (3.2 g, 5.0 mmol) in THF (50 mL) was treated with piperidine (5.0 mL, 50 mmol) and heated to reflux. After 5 hours, the reaction mixture was concentrated and taken up in EtOAc, washed with saturated NaHCO<sub>3</sub>, dried over MgSO<sub>4</sub> and column chromatographed on silica gel using a gradient elution of EtOAc/Hexane to EtOAc. The product (2.7 g) was a white solid with a Mp = 93-95°C; <sup>1</sup>H NMR (DMSO) 7.48-7.46 (m, 4 H), 7.42-7.38 (m, 4 H), 7.38-7.32 (m, 2 H), 7.29 (d, 2 H, J = 8.8 Hz), 7.19 (d, 1 H, J = 9.0 Hz), 7.12-7.10 (m, 3 H), 6.80 (dd, 1 H, J = 8.8, 2.4 Hz), 6.73 (s, 4 H), 5.15 (s, 2 H), 5.13 (s, 2 H), 5.11 (s, 2 H), 3.93 (t, 2

H, J = 5.7 Hz), 2.60-2.50 (m, 2 H), 2.41-2.30 (m, 4 H), 2.15 (s, 3 H), 1.47-1.42 (m, 4 H), 1.36-1.32 (m, 2 H); MS FAB 637 (M+H<sup>+</sup>).

Alternative Procedure

5

Method 6a

Substitution of chlorides

Synthesis illustrated for product No. 76

10 Example No. 76 5-Benzyl-2-(3-benzyl-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole

To a solution of 1.1 g (0.00953 mol) of 5-benzyl-2-(3-benzyl-phenyl)-1-[4-(2-chloro-ethoxy)-benzyl]-3-methyl-1H-indole (example No. 58) in 10 mL of DMF was added 1.1 mL (0.0112 mol) of piperidine, and 0.93 g (0.00561 mol) of potassium iodide. The reaction mixture was heated to ~40-50° C for 4 hours. After cooling the reaction mixture to room temperature, 150 mL of ethyl acetate were added and the mixture was washed with water (3 x 100 mL). The organic solution was collected, washed with saturated brine, removed, dried over magnesium sulfate, filtered and evaporated to yield 1.0 g of product of the product after purification.

Mp = 125 - 126°C; <sup>1</sup>H NMR (DMSO) 7.48 - 7.45 (d, 2 H, J = 7.2 Hz), 7.41 - 7.35 (m, 7 H), 7.33 - 7.28 (m, 2 H), 7.23 - 7.21 (d, 1 H, J = 9.0 Hz), 7.13 - 7.12 (d, 1 H, J = 2.4 Hz), 7.06 - 7.03 (m, 1 H), 6.95 - 6.91 (m, 2 H), 6.83 - 6.80 (dd, 1 H, J = 2.4 Hz, J = 6.3 Hz), 6.75 - 6.70 (m, 4 H), 5.13 (s, 2 H), 5.11 (s, 2 H), 5.02 (s, 2 H), 3.93 - 3.90 (t, 2 H, J = 6.0 Hz), 2.56 - 2.53 (t, 2 H, J = 5.9 Hz), 2.49 - 2.48 (m, 4 H), 2.14 (s, 3 H), 1.46 - 1.40 (m, 4 H), 1.35 - 1.31 (m, 2 H); IR (KBr) 3400, 2900 cm<sup>-1</sup>; MS eI m/z 636 (M+); CHN calcd for C<sub>43</sub>H<sub>44</sub>N<sub>2</sub>O<sub>3</sub> + 0.25 H<sub>2</sub>O.

Physical data for the amine substituted compounds

30 The following compounds were prepared by scheme 13 using method 6. Except for examples No. 76- No. 84 which were prepared using method 6a.

**Example No. 61 5-Benzylxy-2-(4-ethoxy-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole**

Mp = 188 - 191°C;  $^1\text{H}$  NMR (DMSO) 7.45 (d, 2 H, J = 7.3 Hz), 7.40 - 7.25 (m, 5 H), 7.17 (d, 1 H, J = 8.8 Hz), 7.11 (d, 1 H, J = 2.2 Hz), 7.01 (d, 2 H, J = 6.8 Hz), 6.78 (dd, 1 H, J = 8.8 Hz, J = 2.4 Hz), 6.73 (s, 4H), 5.15 (s, 2 H), 5.10 (s, 2H), 4.05 (q, 2H, J = 6.8 Hz), 3.93 (t, 2H, J = 6.0 Hz), 2.55 (t, 2H, J = 5.7 Hz), 2.41 - 2.35 (m, 4 H), 2.14 (s, 3 H), 1.46 - 1.40 (m, 4H), 1.38 - 1.30 (m, 5 H); MS eI m/z 574 (M+).

10. Example No. 62 5-Benzylxy-2-phenyl-3-methyl-1-[4-(2-azepan-1-yl-ethoxy)-benzyl]-1H-indole

Oil;  $^1\text{H}$  NMR (DMSO) 7.50-7.43 (m, 4 H), 7.42-7.37 (m, 5 H), 7.33-7.30(m, 1 H), 7.22 (d, 1 H, J = 8.8 Hz), 7.14 (d, 1 H, J = 2.4 Hz), 6.81 (d, 1 H, J = 6.6 Hz), 6.72 (s, 4 H), 5.18 (s, 2 H), 5.11 (s, 2 H), 3.90 (t, 2 H, J = 6.1 Hz), 2.81-2.75 (m, 2 H), 2.68-2.59 (m, 4 H), 2.16 (s, 3 H), 1.58-1.43 (m, 8 H); MS eI m/z 544 (M+).

**Example No. 64 5-Benzylxy-2-(4-benzylxy-phenyl)-3-methyl-1-[4-(2-azepan-1-yl-ethoxy)-benzyl]-1H-indole**

20. Mp = 106 - 107°C;  $^1\text{H}$  NMR (DMSO) 7.47 (d, 4 H, J = 8.3 Hz), 7.41 - 7.36 (m, 4 H), 7.36 - 7.30 (m, 2 H), 7.29 (d, 2 H, J = 8.8 Hz), 7.19 (d, 1 H, J = 8.8 Hz), 7.14 - 7.10 (m, 3 H), 6.80 (dd, 1 H, J = 8.8 Hz), 6.73 (s, 4 H), 5.15 (s, 2 H), 5.13 (s, 2 H), 5.11 (s, 2 H), 3.90 (t, 2 H, J = 5.9 Hz), 2.76 (t, 2 H, J = 5.9 Hz), 2.64 - 2.56 (m, 4 H), 2.15 (s, 3 H), 1.58 - 1.44 (m, 8 H); MS FAB m/z 651 (M+H+).

25

**Example No. 65 5-Benzylxy-2-(4-benzylxy-phenyl)-3-methyl-1-[4-(2-diisopropylamino-1-yl-ethoxy)-benzyl]-1H-indole**

Mp = 148 - 150°C;  $^1\text{H}$  NMR (DMSO) 7.47 (d, 4 H, J = 8.3 Hz), 7.41 - 7.36 (m, 4 H), 7.36 - 7.32 (m, 2 H), 7.28 (d, 2 H, J = 8.8 Hz), 7.19 (d, 1 H, J = 9.0 Hz), 7.13 - 7.08 (m, 3 H), 6.80 (dd, 1 H, J = 8.8 Hz, 2.4 Hz), 6.76 - 6.68 (m, 4 H), 5.14 (s, 2 H), 5.13 (s, 2 H), 5.11 (s, 2 H), 3.75 (t, 2 H, J = 7.0 Hz), 2.95 (m, 2 H), 2.67 (t, 2 H, J = 7.0 Hz), 2.15 (s, 3 H), 0.93 (d, 12 H, J = 6.4 Hz); MS FAB m/z 653 (M+H+).

Example No. 66 5-Benzyl-2-(4-benzyl-phenyl)-3-methyl-1-[4-(2-butyl-methylamino-1-ylethoxy)-benzyl]-1H-indole

5 Mp = 101 - 104°C;  $^1\text{H}$  NMR (DMSO) 7.45 (d, 4 H,  $J$  = 7.5 Hz), 7.40 - 7.25 (m, 8 H), 7.19 (d, 1 H,  $J$  = 8.8 Hz), 7.12-7.08 (m, 3 H), 6.80 (dd, 1 H,  $J$  = 6.5 Hz,  $J$  = 2.4 Hz), 6.72 (s, 4 H), 5.14 (s, 2 H), 5.13 (s, 2 H), 5.10 (s, 2 H), 3.91 (t, 2 H,  $J$  = 5.9 Hz), 2.64-2.59 (m, 2H), 2.35-2.29 (m, 2 H), 2.17 (s, 3 H), 2.14 (s, 3 H), 1.40-1.31 (m, 2 H), 1.25-1.19 (m, 2 H), 0.83 (t, 3 H, 7.2 Hz); MS esI m/z 638 (M+).

10

Example No. 66a 5-Benzyl-2-(4-benzyl-phenyl)-3-methyl-1-[4-(dimethylamino)-ethoxyl-benzyl]-1H-indole

Mp = 123-124°C

15 Example No. 67 5-Benzyl-2-(4-benzyl-phenyl)-3-methyl-1-[4-[2-(2-methyl-piperidin-1-yl)-ethoxyl-benzyl]-1H-indole

Mp = 121°C

20 Example No. 68 5-Benzyl-2-(4-benzyl-phenyl)-3-methyl-1-[4-[2-(3-methyl-piperidin-1-yl)-ethoxyl-benzyl]-1H-indole

Mp = 90°C

Example No. 69 5-Benzyl-2-(4-benzyl-phenyl)-3-methyl-1-[4-[2-(4-methyl-piperidin-1-yl)-ethoxyl-benzyl]-1H-indole

25 Mp = 98°C;  $^1\text{H}$  NMR (DMSO) 7.46 (d, 4 H,  $J$  = 7.2 Hz), 7.42 - 7.36 (m, 4 H), 7.36 - 7.31 (m, 2 H), 7.28 (d, 2 H,  $J$  = 8.6 Hz), 7.19 (d, 1 H,  $J$  = 9.0 Hz), 7.12 - 7.10 (m, 3 H), 6.80 (dd, 1 H,  $J$  = 8.8 Hz, 2.4 Hz), 6.73 (s, 4 H), 5.15 (s, 2 H), 5.13 (s, 2 H), 5.11 (s, 2 H), 3.93 (t, 2 H,  $J$  = 5.9 Hz), 2.85 - 2.78 (m, 2 H), 2.62 - 2.56 (m, 2 H), 2.15 (s, 3 H), 1.97 - 1.87 (m, 2 H), 1.55 - 1.47 (m, 2 H), 1.30 - 1.20 (m, 1 H), 1.15 - 1.02 (m, 2 H), 0.85 (d, 3 H,  $J$  = 6.6 Hz); MS esI m/z 651 (M+).

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Example No. 70 5-Benzyl-2-(4-benzyloxy-phenyl)-3-methyl-1-[4-[2-((cis)-2,6-Dimethyl-piperidin-1-yl)-ethoxyl-benzyl]-1H-indole  
Mp = 106 - 107°C; <sup>1</sup>H NMR (DMSO) 7.46 (d, 4 H, J = 8.1 Hz), 7.42 - 7.36 (m, 4 H), 7.37 - 7.31 (m, 2 H), 7.29 (d, 2 H, J = 8.8 Hz), 7.18 (d, 1 H, J = 8.8 Hz), 7.14 - 5 7.09 (m, 3 H), 6.80 (dd, 1 H, J = 8.8 Hz, 2.4 Hz), 6.72 (s, 4 H), 5.14 (s, 2 H), 5.13 (s, 2 H), 5.11 (s, 2 H), 3.84 (t, 2 H, J = 7.0 Hz), 2.84 (t, 2 H, J = 6.6 Hz), 2.44 - 2.37 (m, 2 H), 2.15 (s, 3 H), 1.60 - 1.43 (m, 3 H), 1.32 - 1.18 (m, 1 H), 1.16 - 1.06 (m, 2 H), 1.01 (d, 6 H, J = 6.2 Hz).

10 Example No. 71 5-Benzyl-2-(4-benzyloxy-phenyl)-3-methyl-[4-[2-(1,3,3-trimethyl-6-aza-bicyclo[3.2.1]oct-6-yl)-ethoxyl-benzyl]-1H-indole  
Mp = 107°C; MS ESI m/z 705 (M+)+

15 Example No. 71a (1S,4R)-5-Benzyl-2-(4-benzyloxy-phenyl)-3-methyl-[4-[2-(2-Aza-bicyclo [2.2.1] hept-2-yl)-ethoxyl-benzyl]-1H-indole  
The (1S,2R)- 2-aza-bicyclo [2.2.1] heptane used to substitute the bromide was prepared according to the procedure outlined in Syn. Comm. 26(3), 577-584 (1996).  
20 Mp = 95 - 100°C; <sup>1</sup>H NMR (DMSO) 7.32 - 6.55 (m, 21 H), 5.10 - 4.90 (m, 6 H), 3.69 (t, 2 H, J = 5.9 Hz), 2.65 - 2.5 (m, 3 H), 2.10 (s, 2 H), 2.0 (s, 3 H), 1.50 - 1.0 (m, 7 H).

25 Example No. 72 5-Benzyl-2-(4-fluoro-phenyl)-3-methyl-1-[4-(2-azepan-1-yl-ethoxy)-benzyl]-1H-indole  
Oil; <sup>1</sup>H NMR (DMSO) 7.50 - 7.43 (m, 2 H), 7.42 - 7.33 (m, 4 H), 7.32 - 7.20 (m, 4 H), 7.13 (d, 1 H, J = 2.4 Hz), 6.83 (dd, 1 H, J = 2.4 Hz, 6.7 Hz), 6.71 (s, 4 H), 5.14 (s, 2 H), 5.11 (s, 2 H), 3.89 (t, 2 H, J = 5.9 Hz), 3.20 (m, 4 H), 2.74 (t, 2 H, J = 6.0 Hz), 2.15 (s, 3 H), 1.60 - 1.40 (m, 8 H); MS eI m/z 562 (M+).

Example No. 72a 5-Benzylxy-2-(4-fluoro-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole

Oil;  $^1\text{H}$  NMR (DMSO) 7.32 - 6.53 (m, 16 H), 5.00 (s, 2 H), 4.96 (s, 2 H), 3.77 (t, 2 H,  $J$  = 5.8 Hz), 3.22 - 3.14 (m, 4 H), 2.40 (t, 2 H,  $J$  = 5.8 Hz), 2.0 (s, 3 H), 1.29 - 5 1.17 (m, 6 H).

Example No. 72b 5-Benzylxy-2-(4-chloro-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole

Oil;  $^1\text{H}$  NMR (DMSO) 7.52 (d, 2 H,  $J$  = 8.6 Hz), 7.46 (d, 2 H,  $J$  = 6.8 Hz), 7.41 - 10 7.37 (m, 4 H), 7.35 - 7.29 (m, 1 H), 7.25 (d, 1 H,  $J$  = 9.0 Hz), 7.14 (d, 1 H,  $J$  = 2.4 Hz), 6.83 (dd, 1 H,  $J$  = 8.8 Hz, 2.5 Hz), 6.72 - 6.65 (m, 4 H), 5.16 (s, 2 H), 5.11 (s, 2 H), 3.90 (t, 2 H,  $J$  = 5.9 Hz), 2.55 (t, 2 H,  $J$  = 6.0 Hz), 2.41 - 2.26 (m, 4 H), 2.16 (s, 3 H), 1.44 - 1.39 (m, 4 H), 1.38 - 1.29 (m, 2 H); MS eI m/z 564 (M $^+$ ).

15 Example No. 73 5-Benzylxy-2-[3,4-methylenedioxo-phenyl]-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole

Foam;  $^1\text{H}$  NMR (DMSO) 7.45 (d, 2 H,  $J$  = 7.0 Hz), 7.41-7.37 (m, 2 H), 7.33-7.29 (m, 1 H), 7.19 (d, 1 H,  $J$  = 8.8 Hz), 7.11 (d, 1 H,  $J$  = 2.2 Hz), 7.00 (d, 1 H,  $J$  = 7.9 Hz), 6.90 (d, 1 H, 1.4 Hz), 6.82 - 6.78 (m, 2H), 6.74 (s, 4 H), 6.07 (s, 2 H), 5.16 (s, 2 H), 5.10 (s, 2 H), 3.93 (t, 2H,  $J$  = 6.0 Hz), 2.56 (t, 2 H,  $J$  = 6.0Hz), 2.41-2.35 (m, 4H), 2.15 (s, 3H), 1.48-1.41 (m, 4H), 1.38-1.28 (m, 2H); MS eI m/z 574 (M $^+$ ).

25 Example No. 74 5-Benzylxy-2-[4-isopropoxy-phenyl]-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole

Foam;  $^1\text{H}$  NMR (DMSO) 7.46 (d, 2H,  $J$  = 7.7 Hz), 7.42 - 7.28 (m, 3 H), 7.25 (d, 2 H,  $J$  = 8.7 Hz), 7.17 (d, 1 H,  $J$  = 8.7 Hz), 7.11 (d, 1 H,  $J$  = 2.4 Hz), 6.99 (d, 2 H,  $J$  = 8.6 Hz), 6.79 (dd, 1 H,  $J$  = 2.4 Hz, 8.8 Hz), 6.73 (s, 4 H), 5.14 (s, 2 H), 5.10 (s, 2 H), 4.70 - 4.60 (m, 1 H), 3.92 (t, 2 H,  $J$  = 5.7 Hz), 2.55 (t, 2 H, 5.7 Hz), 2.40 - 2.30 (bs, 4 H), 2.15 (s, 3 H), 1.50 - 1.40 (m, 4 H), 1.40 - 30 1.30 (m, 2 H), 1.28 (d, 6 H,  $J$  = 6.2 Hz); MS eI m/z 588 (M $^+$ ).

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Example No. 75 5-Benzyl oxy-2-[4-methyl-phenyl]-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole

Oil;  $^1\text{H}$  NMR (DMSO) 7.46 (d, 2 H,  $J = 7.2$  Hz), 7.45 - 7.18 (m, 8 H), 7.12 (d, 1 H,  $J = 2.4$  Hz), 6.81 (dd, 1 H,  $J = 2.4$  Hz, 8.6 Hz), 6.73 (s, 4 H), 5.15 (s, 2 H), 5.10 (s, 2 H), 3.92 (t, 2 H,  $J = 5.9$  Hz), 2.55 (t, 2 H,  $J = 5.9$  Hz), 2.45 - 2.30 (m, 7 H), 2.10 (s, 3 H), 1.50 - 1.40 (m, 4 H), 1.48 - 1.35 (m, 2 H); MS eI m/z 544 (M+).

Example No. 77 1-[4-(2-Azepan-1-yl-ethoxy)-benzyl]-5-benzyl oxy-2-(3-benzyl oxy-phenyl)-3-methyl-1H-indole  
10  $M_p = 103 - 105^\circ\text{C}$ ;  $^1\text{H}$  NMR (DMSO) 7.47 - 7.45 (d, 2 H,  $J = 8.1$  Hz), 7.41 - 7.35 (m, 7 H), 7.32 - 7.29 (t, 2 H, 7.0 Hz), 7.23 - 7.21 (d, 1 H,  $J = 8.7$  Hz), 7.13 - 7.12 (d, 1 H,  $J = 2.1$  Hz), 7.06 - 7.03 (m, 1 H), 6.95 - 6.91 (m, 2 H), 6.83 - 6.80 (m, 1 H), 6.75 - 6.73 (m, 4 H), 5.13 (s, 2 H), 5.11 (s, 2 H), 5.02 (s, 2 H), 3.90 - 3.87 (t, 2 H,  $J = 6.0$  Hz), 2.76 - 2.73 (t, 2 H,  $J = 6.0$  Hz), 2.49 - 2.48 (m, 4 H), 2.13 (s, 3 H), 1.51 (s, 8 H); IR 3400, 2900  $\text{cm}^{-1}$ ; MS eI m/z 650 (M+); CHN calcd for  $\text{C}_{44}\text{H}_{46}\text{N}_2\text{O}_3$ .

Example No. 78 5-Benzyl oxy-2-(4-benzyl oxy-3-fluoro-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole  
20  $M_p = 125-128^\circ\text{C}$ ;  $^1\text{H}$  NMR (DMSO) 7.50 - 7.45 (m, 4 H), 7.43 - 7.28 (m, 7 H), 7.26 - 7.20 (m, 2 H), 7.14 - 7.09 (m, 2 H), 6.82 (dd, 1 H,  $J = 2.4$  Hz, 8.8 Hz), 6.72 (s, 4 H), 5.21 (s, 2 H), 5.16 (s, 2 H), 5.11 (s, 2 H), 3.94 (t, 2 H,  $J = 5.8$  Hz), 2.62 - 2.56 (m, 2 H), 2.41 - 2.36 (m, 4 H), 2.15 (s, 3 H), 1.45 - 1.40 (m, 4 H), 1.40 - 1.31 (m, 2 H); MS eI m/Z 654 (M+); CHN calcd for  $\text{C}_{43}\text{H}_{43}\text{FN}_2\text{O}_3$ .

Example No. 79 5-Benzyl oxy-2-(4-benzyl oxy-3-fluoro-phenyl)-3-methyl-1-[4-(2-azepan-1-yl-ethoxy)-benzyl]-1H-indole  
30  $M_p = 122-124^\circ\text{C}$ ;  $^1\text{H}$  NMR (DMSO) 7.50 - 7.28 (m, 10 H), 7.26 - 7.20 (m, 2 H), 7.15 - 7.10 (m, 2 H), 6.88 - 6.76 (m, 2 H), 6.70 (s, 4 H), 5.22 (s, 2 H), 5.16 (s, 2 H), 5.11 (s, 2 H), 3.92 - 3.86 (m, 2 H), 2.82 - 2.65 (m, 2 H), 2.65 - 2.55 (m, 4 H), 2.15 (s, 3 H), 1.60 - 1.4 (m, 8 H); MS eI m/Z 668 (M+); CHN calcd for  $\text{C}_{44}\text{H}_{45}\text{FN}_2\text{O}_3$ .

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Example No. 80 5-Benzyl oxy-2-(3-methoxy-phenyl)-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-3-methyl-1H-indole

Mp 86 - 87°C; <sup>1</sup>H NMR (DMSO) 7.50 - 7.49 (m, 2 H), 7.46 - 7.31 (m, 4 H), 7.24 -

5 7.21 (d, 1 H, J = 8.8 Hz), 7.15 - 7.14 (d, 1 H, J = 2.3 Hz), 7.00 - 6.93 (m, 2 H),  
6.88 - 6.81 (m, 2 H), 6.75 (s, 4 H), 5.18 (s, 2 H), 5.12 (s, 2 H), 3.96 - 3.92  
(t, 2 H, J = 5.9 Hz), 3.71 (s, 3 H), 2.59 - 2.55 (t, 2 H, J = 5.8 Hz), 2.37 (s, 4 H),  
2.18 (s, 3 H), 1.49 - 1.42 (m, 4 H), 1.37 - 1.34 (m, 2 H); MS eI m/z 561 (M+); CHN  
calcd for C<sub>37</sub>H<sub>40</sub>N<sub>2</sub>O<sub>3</sub> + 0.25 H<sub>2</sub>O.

10

Example No. 81 5-Benzyl oxy-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-2-(4-trifluoromethoxy-phenyl)-1H-indole

Mp = 107 - 108°C; <sup>1</sup>H NMR (DMSO) 7.52 - 7.45 (m, 6 H), 7.41 - 7.26 (m, 4 H),

15 7.17 - 7.16 (d, 1 H, J = 2.3 Hz), 6.87 - 6.84 (dd, 1 H, J = 2.3 Hz, J = 6.4 Hz), 6.75  
- 6.68 (m, 4 H), 5.18 (s, 2 H), 5.13 (s, 2 H), 3.95 - 3.91 (t, 2 H, J = 5.9 Hz), 2.58 -  
2.54 (t, 2 H, J = 5.9 Hz), 2.38 - 2.34 (m, 4 H), 2.17 - 2.15 (s, 3 H), 1.49 - 1.42  
(m, 4 H), 1.35 - 1.34 (d, 2 H, J = 4.9 Hz); IR 3400, 2900, 1600 cm<sup>-1</sup>; MS eI m/z 615  
(M+); CHN calcd for C<sub>37</sub>H<sub>37</sub>F<sub>3</sub>N<sub>2</sub>O<sub>3</sub>.

20

Example No. 82 (2-[4-[5-Benzyl oxy-2-(4-benzyl oxy-phenyl)-3-methyl-indol-1-ylmethyl]-phenoxy]-ethyl)-cyclohexyl-amine

Mp = 87-90°C; <sup>1</sup>H NMR (DMSO) 7.46 (dd, 4H, J = 6.9Hz, 0.6Hz), 7.42-7.27

(m, 9H), 7.19 (d, 1H, J = 9Hz), 7.14-7.08 (m, 3H), 6.80 (dd, 1H, J = 6.4Hz, 2.4Hz),  
25 6.75- 6.70 (m, 4H), 5.15(s, 2H), 5.13 (s, 2H), 5.13(s, 2H), 3.89 (t, 2H, J = 5.6),  
2.84 (m, 2H), 2.48 (m, 1H), 2.14 (s, 3H), 1.80 ( m, 2H), 1.65 ( m, 2H), 1.61  
(m, 1H), 0.96-1.19 (m, 5H); MS eI m/Z 650 (M+); CHN calcd for C<sub>44</sub>H<sub>46</sub>N<sub>2</sub>O<sub>4</sub>.

30

Example No. 83 5-Benzyl oxy-2-(4-benzyl oxy-phenyl)-3-methyl-1-[4-methylpiperazin-1-yl]-ethoxyl-benzyl]-1H-indole

Mp = 88-91°C; <sup>1</sup>H NMR (DMSO) 7.47 (m, 4H), 7.26-7.42 (m, 8H), 7.19

(d, 1H, J = 8.8), 7.10-1.12 (m, 3H), 6.80 (q, 1H, J = 6.3Hz, 2.4Hz), 6.73 (m, 4H),  
5.15 (s, 2H), 5.13 (s, 2H), 5.11 (s, 2H), 3.94 (t, 2H, J = 5.9Hz), 2.59 (t, 2H), 2.42

(m, 4H), 2.29 (m, 4H), 2.15 (s, 3H), 2.12 (s, 3H); MS eI m/Z 652 (M+); CHN calcd for  $C_{43}H_{45}N_3O_3$ .

Example No. 84 1-[4-(2-Azepan-1-yl-ethoxy)-benzyl]-5-benzyl oxy-2-(3-methoxy-phenyl)-3-methyl-1H-indole  
 5  $M_p = 103 - 105^{\circ}\text{C}$ ;  $^1\text{H}$  NMR (DMSO) 7.47 - 7.45 (d, 2 H,  $J = 8.1$  Hz), 7.41 - 7.35 (m, 7 H), 7.32 - 7.29 (t, 2 H, 7.0 Hz), 7.23 - 7.21 (d, 1 H,  $J = 8.7$  Hz), 7.13 - 7.12 (d, 1 H,  $J = 2.1$  Hz), 7.06 - 7.03 (m, 1 H), 6.95 - 6.91 (m, 2 H), 6.83 - 6.80 (m, 1 H), 6.75 - 6.73 (m, 4 H), 5.13 (s, 2 H), 5.11 (s, 2 H), 5.02 (s, 2 H), 3.90 - 3.87 (t, 2 H,  $J = 6.0$  Hz), 2.76 - 2.73 (t, 2 H,  $J = 6.0$  Hz), 2.49 - 2.48 (m, 4 H), 2.13 (s, 3 H), 1.51 (s, 8 H); IR 3400, 2900  $\text{cm}^{-1}$ ; MS eI m/z 650 (M+); CHN calcd for  $C_{44}H_{46}N_2O_3$ .

Data and procedures for compounds From Table 11 (ER Receptor Data Table, *infra*) of Text

15

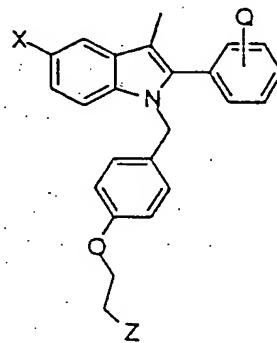
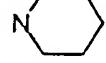
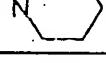
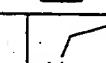
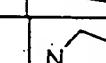
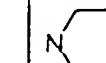
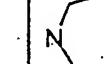


Table 7

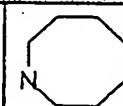
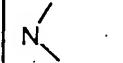
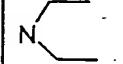
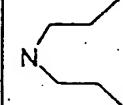
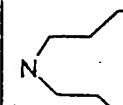
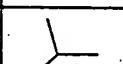
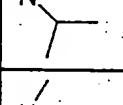
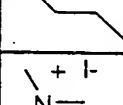
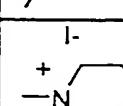
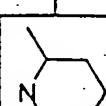
Example No.	X	Q	Z
No. 85	H	H	
No. 86	H	4'-OH	

Table 7 (Cont'd)

Example No.	X	Q	Z
No. 87	OH	H	
No. 88	OMe	4'-OH	
No. 89	OH	4'-OMe	
No. 90	OMe	4'-OMe	
No. 91	OMe	4'-OMe	
No. 92	OH	4'-OEt	
No. 93	OH	4'-OEt	
No. 94	F	4'-OH	
No. 95	OH	H	
No. 96	OH	4'-OH	
No. 97	OH	4'-OH	
No. 98	OH	4'-OH	

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Table 7 (Cont'd)

Example No.	X	Q	Z
No. 99	OH	4'-OH	
No. 100	OH	4'-OH	
No. 101	OH	4'-OH	
No. 102	OH	4'-OH	
No. 103	OH	4'-OH	
No. 104	OH	4'-OH	
No. 105	OH	4'-OH	
No. 106	OH	4'-OH	
No. 107	OH	4'-OH	
No. 108	OH	4'-OH	

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Table 7 (Cont'd)

Example No.	X	Q	Z
No. 109	OH	4'-OH	
No. 110	OH	4'-OH	
No. 111	OH	4'-OH	
No. 112	OH	4'-OH	
No. 113	OH	4'-OH	
No. 114	OH	4'-OH	
No. 115	OH	4'-OH	
No. 116	OH	4'-F	
No. 117	OH	4'-F	
No. 118	OH	3'-OMe,4'-OH	
No. 119	OH	3',4'-OCH <sub>2</sub> O-	

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Table 7 (Cont'd)

Example No.	X	Q	Z
No. 120	OH	4'-O-iPr	
No. 121	OH	4'-O-iPr	
No. 122	OH	4'-O-Cp	
No. 123	OH	4'-CF <sub>3</sub>	
No. 124	OH	4'-CH <sub>3</sub>	
No. 125	OH	4'-Cl	
No. 126	OH	2',4',-Dimethoxy	
No. 127	OH	3'-OH	
No. 128	OH	3'-OH	
No. 129	OH	4'-OH,3'-F	
No. 130	OH	4'-OH, 3'-F	
No. 131	OH	3'-OMe	
No. 132	OH	4'-OCF <sub>3</sub>	

Hydrogenation of Indoles Containing Benzyl Ether(s)

Method 7  
Illustrated For Example No. 97

5    2-(4-Hydroxy-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-  
1H-indol-5-ol

A suspension of 10% Pd/C (1.1 g) in EtOH was treated with a solution of No. 63 (2.2 g, 3.4 mmol) in THF/EtOH. Cyclohexadiene (6.0 mL, 63 mmol) was added 10 and the reaction was stirred for 48 hours. The catalyst was filtered through Celite and the reaction mixture was concentrated and chromatographed on silica gel using a gradient elution of MeOH/CH<sub>2</sub>Cl<sub>2</sub> (1:19 to 1:10) to yield 0.8 g of the product as a white solid. Mp = 109-113°C; CHN calc'd for C<sub>29</sub>H<sub>32</sub>N<sub>2</sub>O<sub>3</sub> + 0.5 H<sub>2</sub>O; <sup>1</sup>H NMR 9.64 (s, 1 H), 8.67 (s, 1 H), 7.14 (d, 2 H, J = 8.6 Hz), 7.05 (d, 1 H, J = 8.6 Hz), 6.84 (d, 2 H, 15 J = 8.8 Hz), 6.79 (d, 1 H, J = 2.2 Hz), 6.74 (s, 4 H), 6.56 (dd, 1 H, J = 8.8, 2.4 Hz), 5.09 (s, 2 H); 3.95-3.93 (m, 2 H), 2.60-2.51 (m, 2 H), 2.39-2.38 (m, 4 H), 2.09 (s, 3 H), 1.46-1.45 (m, 4 H), 1.35-1.34 (m, 2 H); IR (KBr) 3350 (br), 2920, 1620, 1510 cm<sup>-1</sup>; MS (EI) m/z 456.

20    Alternatively, the compounds may be dissolved in a THF/EtOH solution (or other appropriate solvent) and hydrogenated with H<sub>2</sub> and 10% Pd/C using either a ballon or Parr Hydrogenator. Either procedure is effective. In many of the examples, the compounds were made into acid addition salts. The procedure for the preparation of an HCl salt is given below (Method 8).

25

**Method 8**

1.0 g of Example No. 97 free base from the hydrogenation procedure above in a large test tube was dissolved in 20 mL of MeOH. This was treated with slow addition of 2.6 mL 1.0 N HCl and then 4.0 mL deionized water. The tube was 30 partially opened to the atmosphere to encourage slow evaporation of the solvents. After about ten minutes, crystals began to appear and after 4 hours the solution was filtered and the solid crystals washed with water. The product was present as 0.42 g of white crystalline plates with a melting point of 184-185°C. The mother liquor yielded an

additional crop of 0.30 g of white solid with a melting point of 177-182°C. CHN calc'd for  $C_{29}H_{32}N_2O_3 + HCl + 1 H_2O$ .

Alternatively, the compounds can be made into quaternary ammonium salts. An 5 example procedure for the synthesis of example No. 107 is given below (Method 9).

### Method 9

10 Example No. 107 2-(4-Hydroxy-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl-1H-indol-5-ol methiodide

0.8g of example No. 97 was dissolved in 18 mL THF and treated with 2 mL of methyl iodide. The solution was heated to reflux for an hour. The reaction was allowed to come to room temperature and the solids filtered to yield 0.72 g as a crystalline solid.

Mp = 214 - 217°C, CHN calcd for  $C_{29}H_{32}N_2O_3 + CH_3I + 0.5 H_2O$ .

15

Example No. 106 2-(4-Hydroxy-phenyl)-3-methyl-1-[4-(2-dimethyl-1-yl-ethoxy)-benzyl-1H-indol-5-ol methiodide was prepared similarly to No.

106 except using No. 100 for starting material: Mp = 245 - 250°C;  $^1H$  NMR (DMSO)

9.66 (s, 1 H), 8.69 (s, 1 H), 7.16 (d, 2 H,  $J = 8.4$  Hz), 7.05 (d, 1 H,  $J = 8.8$  Hz),

20 6.84 (d, 1 H,  $J = 8.6$  Hz), 6.81 - 6.75 (m, 6H), 6.56 (dd, 1 H,  $J = 2.4$  Hz, 8.7 Hz),

5.12 (s, 2 H), 4.34 (m, 2 H), 3.70 (t, 2 H,  $J = 4.6$  Hz), 3.11 (s, 9 H), 2.09 (s, 3 H);

IR (KBr) 3250, 1500, 1250; MS eI m/z 416 (M+); CHN calcd for  $C_{26}H_{28}N_2O_3 + 1.09 CH_3I + 0.8 H_2O$ .

25

### Physical Data for final, deprotected compounds

The following compounds are either free bases, HCl salts or acetate salts. They were prepared according to the procedure outlined in method 7 using the appropriate benzyl ether for precursor. Where a compound from table 1 does not contain a free 30 phenolic functionality, then it was unnecessary to debenzylate it and method 7 not applied. The physical data for these compounds (No. 85, No. 90- No. 91) is still presented below.

**Example No. 85 4-{3-Methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole} (HCl)**

5 Mp = 134 - 137°C; <sup>1</sup>H NMR (DMSO) 10.33 (s, 1H), 7.56 - 7.38 (m, 6 H), 7.32 (d, 1 H, J = 8.1 Hz), 7.14 - 7.0 (m, 2 H), 6.80 (s, 4 H), 5.24 (s, 2 H), 4.28 (t, 2 H, J = 5.0 Hz), 3.50 - 3.40 (m, 4 H), 3.0 - 2.95 (m, 2 H), 2.10 (s, 3 H), 1.80 - 1.60 (m, 5 H), 1.40 - 1.35 (m, 1 H); IR 3400, 2900, 1510, 1250 cm<sup>-1</sup>; MS (+) FAB m/z 425 [M+H]<sup>+</sup>; CHN calcd for C<sub>29</sub>H<sub>32</sub>N<sub>2</sub>O + 1.0 HCl + 1.0 H<sub>2</sub>O.

10 **Example No. 86 4-{3-Methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indol-2-yl}-phenol hydrochloride (HCl)**

15 Mp = 192 - 194°C; <sup>1</sup>H NMR (DMSO), 10.28 (s, 1 H), 9.75 (s, 1 H), 7.51 - 7.49 (m, 1H), 7.27 (dd, 1 H, J = 7.0 Hz, 0.7 Hz), 7.18 (d, 2 H, J = 7.6 Hz), 7.09 - 7.02 (m, 2 H), 6.86 (d, 2 H, J = 8.6 Hz), 6.80 (s, 4 H), 5.20 (s, 2 H), 4.28 (t, 2 H, J = 4.9 Hz), 3.50 - 3.35 (m, 4 H), 3.0 - 2.85 (m, 2 H), 2.20 (s, 3 H), 1.80 - 1.60 (m, 5 H), 1.40 - 1.30 (m, 1 H); IR 3400, 3100, 2600, 1500, 1225 cm<sup>-1</sup>; MS eI m/z 440 (M<sup>+</sup>); CHN calc for C<sub>29</sub>H<sub>32</sub>N<sub>2</sub>O<sub>2</sub> + 1 HCl.

20 **Example No. 87 3-Methyl-2-phenyl-1-[4-(2-piperidine-1-yl-ethoxy)-benzyl]-1H-indol-5-ol (HCl)**

25 Mp = 228-230°C; <sup>1</sup>H NMR 10.1 (brs, 1 H), 8.76 (s, 1 H), 7.55 - 7.45 (m, 5 H), 7.10 (d, 1 H, J = 8.8 Hz), 6.85 - 6.80 (m, 5 H), 6.61 (d, 1 H, J = 8.8 Hz), 5.15 (s, 2 H), 4.25 (t, 2 H, J = 4.8 Hz), 3.47-3.35 (m, 4 H), 2.96-2.87 (m, 2 H), 2.12 (s, 3 H), 1.75-1.65 (m, 5 H), 1.31-1.28 (m, 1 H); MS eI m/z 440 (M<sup>+</sup>); CHN calcd for C<sub>29</sub>H<sub>32</sub>N<sub>2</sub>O<sub>2</sub> + 1 HCl + .33 H<sub>2</sub>O; IR (KBr) 3200, 2500, 1450, 1200 cm<sup>-1</sup>.

30 **Example No. 88 4-{5-Methoxy-3-methyl-1-[4-[2-(piperidin-1-yl)-ethoxyl-benzyl]-1H-indol-2-yl}-phenol**

Mpt = 87-90°C; <sup>1</sup>H NMR (DMSO) 9.67 (s, 1 H), 7.16 (d, 2 H, J = 8.6 Hz), 7.16 (1 H buried), 6.98 (d, 1 H, J = 2.4 Hz), 6.85 (d, 2 H, J = 8.6 Hz), 6.73 (s, 4 H), 6.69 (dd, 1 H, J = 8.8, 2.4 Hz), 5.13 (s, 2 H), 3.94 (t, 2 H, J = 5.7 Hz), 3.76 (s, 3 H), 2.63-2.50 (m, 2 H), 2.43-2.31 (m, 4 H), 2.15 (s, 3 H), 1.49-1.40 (m, 4 H), 1.39-

1.25 (m, 2 H); IR (KBr) 3400 (br), 2920, 1610, 1520  $\text{cm}^{-1}$ ; MS eI m/z 470; CHN  
calcd for  $\text{C}_{30}\text{H}_{34}\text{N}_2\text{O}_3 + 0.1 \text{ H}_2\text{O}$ .

Example No. 89 2-(4-methoxy-phenyl)-3-methyl-1-[4-[2-(piperidin-1-yl)-ethoxy]-benzyl]-1H-indol-5-ol  
5 Mp = 188-189°C;  $^1\text{H}$  NMR (DMSO) 8.70  
(s, 1 H), 7.27 (d, 2 H, J = 8.6 Hz), 7.06 (d, 1 H, J = 8.6 Hz), 7.02 (d, 2 H, J = 8.8 Hz), 6.81 (d, 1 H, J = 2.2 Hz), 6.73 (s, 4 H), 6.58 (dd, 1 H, J = 8.8, 2.4 Hz), 5.10 (s, 2 H), 3.93 (t, 2 H, J = 5.9 Hz), 3.79 (s, 3 H), 2.56 (t, 2 H, J = 5.9 Hz), 2.41-10 2.32 (m, 4 H), 2.10 (s, 3 H), 1.47-1.41 (m, 4 H), 1.34-1.31 (m, 2 H); MS eI m/z 470; CHN calcd for  $\text{C}_{30}\text{H}_{34}\text{N}_2\text{O}_3 + 0.1 \text{ H}_2\text{O}$ .

Example No. 90 5-Methoxy-2-(4-methoxy-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole (HCl)  
15 Mp = 188-191°C;  $^1\text{H}$  NMR (DMSO) 10.35 (brs, 1 H), 7.27 (d, 2 H, J = 8.8 Hz), 7.17 (d, 1 H, J = 8.8 Hz), 7.03 (d, 2 H, J = 8.6 Hz), 6.99 (d, 1 H, J = 2.5 Hz), 6.82 - 6.78 (m, 4 H), 6.71 (dd, 1 H, J = 8.8 Hz, J = 2.5 Hz), 5.17 (s, 2 H), 4.31-4.22 (m, 2 H), 3.79 (s, 3 H), 3.76 (s, 3 H), 3.43 - 3.36 (m, 4 H), 2.97 - 2.83 (m, 2 H), 2.16 (s, 3 H), 1.80 - 1.59 (m, 5 H), 1.41 - 1.26 (m, 1 H); IR (KBr) 2920, 1450, 1250  $\text{cm}^{-1}$ ; MS eI m/z 484 (M+); CHN calc for  $\text{C}_{31}\text{H}_{36}\text{N}_2\text{O}_3 + 1 \text{ HCl}$ .

Example No. 91 1-[4-(2-Azepan-1-yl-ethoxy)-benzyl]-5-methoxy-2-(4-methoxy-phenyl)-3-methyl-1H-indole (HCl)  
Mp = 161-163°C;  $^1\text{H}$  NMR (DMSO) 10.65 (brs, 1 H), 7.27  
25 (d, 2 H, J = 8.8 Hz), 7.17 (d, 1 H, J = 8.8 Hz), 7.03 (d, 2 H, J = 8.6 Hz), 6.99 (d, 1 H, J = 2.5 Hz), 6.82 - 6.77 (m, 4 H), 6.71 (dd, 1 H, J = 8.8 Hz, J = 2.5 Hz), 5.17 (s, 2 H), 4.27 (m, 2 H), 3.79 (s, 3 H), 3.76 (s, 3 H), 3.44 - 3.30 (m, 4 H), 3.17 (m, 2 H), 2.16 (s, 3 H), 1.82 - 1.77 (m, 4 H), 1.63 - 1.48 (m, 4 H); MS eI m/z 499 (M+); CHN calc for  $\text{C}_{32}\text{H}_{38}\text{N}_2\text{O}_3 + 1 \text{ HCl}$ .

**Example No. 92 2-(4-Ethoxy-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indol-5-ol**

Mp = 173-175°C;  $^1\text{H}$  NMR (DMSO) 8.69 (s, 1 H), 7.25 (d, 2 H, J = 8.8Hz), 7.04 (d, 1H, J = 8.8 Hz), 6.99 (dd, 2 H, J = 6.8 Hz, J = 2.0 Hz),  
5 6.80 (d, 1 H, J = 2.2 Hz), 6.73 (s, 4H), 6.59 (dd, 1 H, J = 8.5 J = 2.2),  
5.09 (s, 2H), 4.05 (q, 2 H, J = 7.03 Hz), 3.93 (t, 2 H, J = 6.0 Hz), 2.62 - 2.56 (m, 2H), 2.41 - 2.36 (m, 4 H), 2.09 (s, 3H), 1.45 - 1.41 (m, 4H), 1.38 - 1.30 (m, 5H); MS eI m/z 484 (M+); CHN calc for  $\text{C}_{31}\text{H}_{38}\text{N}_2\text{O}_3$  + .25H<sub>2</sub>O.

10 Example No. 93 1-[4-(2-Azepan-1-yl-ethoxy)-benzyl]-2-(4-ethoxy-phenyl)-3-methyl-1H-indol-5-ol

Mp = 133-135°C;  $^1\text{H}$  NMR (DMSO) 8.69 (s, 1 H), 7.25 (d, 2 H, J = 8.8Hz), 7.04 (d, 1H, J = 8.8 Hz), 6.99 (dd, 2 H, J = 6.8 Hz, J = 2.0 Hz),  
6.80 (d, 1 H, J = 2.2Hz), 6.73 (s, 4H), 6.59 (dd, 1 H, J = 8.5 Hz, J = 2.2 Hz),  
15 5.09 (s, 2H), 4.05 (q, 2H, J = 7.03 Hz), 3.90 (t, 2H, J = 6.1 Hz), 2.75 (t, 2H, J = 6.0 Hz), 2.62 - 2.58 (m, 4 H), 2.09 (s, 3 H), 1.58 - 1.44 (m, 8 H), 1.33 (t, 3H, J = 7.0 Hz); IR (KBr) 2930, 1470, 1250 CM<sup>-1</sup>; MS eI m/z 498 (M+); CHN calc for  $\text{C}_{32}\text{H}_{38}\text{N}_2\text{O}_3$ .

20 Example No. 94 4-[5-Fluoro-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indol-2-yl]-phenol (HCl)

Mp = 223-225°C;  $^1\text{H}$  NMR (DMSO) 10.30 (br s, 1H), 7.27 - 7.23 (m, 2 H),  
7.17 (d, 2 H, J = 8.6 Hz), 6.88 - 6.79 (m, 7H), 5.20 (s, 2H), 4.28 (t, 2H, J = 5.0 Hz), 3.42 - 3.35 (m, 4 H), 3.00 - 2.85 (m, 2 H), 2.14 (s, 3 H),  
25 1.78 - 1.70 (m, 4 H), 1.67 - 1.59 (m, 1 H), 1.40 - 1.26 (m, 1 H); MS eI m/z 458 (M+).

**Example No. 95 1-[4-(2-Azepan-1-yl-ethoxy)-benzyl]-3-methyl-2-phenyl-1H-indol-5-ol (HCl)**

30 Mp = 203-204°C;  $^1\text{H}$  NMR (DMSO) 10.50 (brs, 1 H), 8.80 (s, 1 H), 7.50 - 7.38 (m, 5 H); 7.10 (d, 1 H, J = 8.8Hz), 6.83 - 6.77 (m, 5 H), 6.60 (d, 1 H, J = 6.6 Hz), 5.15 (s, 2H), 4.26 (t, 2 H, J = 5.2 Hz), 3.45 - 3.35 (m, 4 H), 3.21-3.10 (m,

2 H), 2.12 (s, 3H), 1.85-1.75 (m, 4 H), 1.70 - 1.51 (m, 4 H); MS eI m/z 454 (M+); CHN calc for  $C_{30}H_{34}N_2O_2$  + 1 HCl.

Example No. 96 2-(4-Hydroxy-phenyl)-3-methyl-1-[4-(2-pyrollidin-1-yl-ethoxy)-benzyl]-1H-indol-5-ol

5 Mp = 105-110°C; CHN calc'd for  $C_{28}H_{30}N_2O_3$  + 0.4 H<sub>2</sub>O; <sup>1</sup>H NMR (DMSO) 9.65 (s, 1 H), 8.67 (s, 1 H), 7.15 (d, 2 H, J = 8.6 Hz), 7.05 (d, 1 H, J = 8.6 Hz), 6.84 (d, 2 H, J = 2 H), 6.79 (d, 1 H, J = 2.4 Hz), 6.56 (dd, 1 H, J = 8.6, 2.2 Hz), 6.74 (s, 4 H), 5.09 (s, 2 H), 3.95 (t, 2 H, J = 5.7 Hz), 3.39-3.23 (m, 4 H), 2.80-2.75 (m, 2 H),  
10 2.09 (s, 3 H), 1.67-1.64 (m, 4 H); IR (KBr) 3410 (br), 1620, 1510 cm<sup>-1</sup>; MS (EI) m/z 442

Example No. 98 1-[4-(2-Azepan-1-yl-ethoxy)-benzyl]-2-(4-hydroxy-phenyl)-3-methyl-1H-indol-5-ol (HCl)

15 Mp = 168 - 171°C; <sup>1</sup>H NMR (DMSO) 10.11 (br s, 1 H), 9.70 (s, 1 H), 8.71 (s, 1 H); 7.15 (d, 2 H, J = 8.6 Hz), 7.05 (d, 1 H, J = 8.8 Hz), 6.85 (d, 2 H, J = 8.8 Hz), 6.80 - 6.77 (m, 5 H), 6.56 (dd, 1 H, J = 8.8 Hz, 2.2 Hz), 5.11 (s, 2 H), 4.26 (t, 2 H, J = 4.6 Hz), 3.48 - 3.30 (m, 4 H), 3.22 - 3.08 (m, 2 H), 2.09 (s, 3 H), 1.83 - 1.76 (m, 4 H), 1.67 - 1.48 (m, 4 H); IR (KBr) 3500 br, 3250 br, 2900, 1610; MS FAB m/z 471 (M+H+); CHN calcd for  $C_{30}H_{34}N_2O_3$  + 2.5 H<sub>2</sub>O + HCl.

Example No. 98 Acetate Salt

Made by the precipitation of No. 98 free base from acetone and acetic acid.

Mp = 174 - 178°C

25

Example No. 99 1-[4-(2-Azocan-1-yl-ethoxy)-benzyl]-2-(4-hydroxy-phenyl)-3-methyl-1H-indol-5-ol

Mp = 98 - 102°C; <sup>1</sup>H NMR (DMSO) 9.63 (s, 1 H), 8.68 (s, 1 H), 7.15 - 7.13 (m, 2 H), 7.05 (d, 1 H, J = 8.5 Hz), 6.83 (dd, 2 H, J = 2.0 Hz, 6.6 Hz), 6.79 (d, 1 H, J = 2.2 Hz), 6.73 (s, 4 H), 6.55 (dd, 1 H, J = 2.2 Hz, 8.6 Hz), 5.08 (s, 2 H), 3.89 (t, 2 H, J = 5.7 Hz), 2.74 (t, 2 H, J = 5.4 Hz), 2.55 (bs, 4 H), 2.08 (s, 3 H), 1.55 (s, 2

H), 1.46 (s, 8 H); IR 3400, 2900, 1250  $\text{cm}^{-1}$ ; MS eI m/z 484 (M+); CHN calcd for  $\text{C}_{31}\text{H}_{36}\text{N}_2\text{O}_3 + .30 \text{ H}_2\text{O}$ .

5 Example No. 100 2-(4-Hydroxy-phenyl)-3-methyl-1-[4-(2-dimethyl-1-yl-ethoxy)-benzyl]-1H-indol-5-ol

Mp = 95 - 105°C; IR (KBr) 3400 br, 2900, 1610  $\text{cm}^{-1}$ ; MS eI m/z 416 (M+); CHN calcd for  $\text{C}_{26}\text{H}_{23}\text{N}_2\text{O}_3 + 0.5 \text{ H}_2\text{O}$ .

10 Example No. 101 2-(4-Hydroxy-phenyl)-3-methyl-1-[4-(2-diethyl-1-yl-ethoxy)-benzyl]-1H-indol-5-ol

15 Mp = 100-107°C; CHN calc'd for  $\text{C}_{28}\text{H}_{32}\text{N}_2\text{O}_3 + 0.25 \text{ H}_2\text{O}$ ;  $^1\text{H}$  NMR (DMSO) 9.64 (s, 1 H), 8.67 (s, 1 H), 7.14 (d, 2 H,  $J = 8.6 \text{ Hz}$ ), 7.05 (d, 1 H,  $J = 8.8 \text{ Hz}$ ), 6.84 (d, 2 H,  $J = 8.6 \text{ Hz}$ ), 6.79 (d, 1 H, 2.2 Hz), 6.74 (s, 4 H), 6.56 (dd, 1 H,  $J = 8.8, 2.4 \text{ Hz}$ ), 5.09 (s, 2 H), 3.95-3.85 (m, 2 H), 2.80-2.60 (m, 2 H), 2.58-2.40 (m, 4 H), 2.09 (s, 3 H), 0.93 (t, 6 H,  $J = 7.0 \text{ Hz}$ ); IR (KBr) 3410 (br), 2950, 1610, 1510  $\text{cm}^{-1}$ ; MS FAB 445 (M+H+).

20 Example No. 102 1-[4-(2-Dipropylamino-ethoxy)-benzyl]-2-(4-hydroxy-phenyl)-3-methyl-1H-indol-5-ol

25 Mp = 83 - 86°C;  $^1\text{H}$  NMR (DMSO) 9.64 (s, 1 H), 8.67 (s, 1 H), 7.14 (d, 2 H,  $J = 8.6$ ), 7.04 (d, 1 H,  $J = 8.6 \text{ Hz}$ ), 6.83 (d, 2 H,  $J = 8.6 \text{ Hz}$ ), 6.78 (d, 1 H,  $J = 2.2 \text{ Hz}$ ), 6.72 (m, 4 H), 6.55 (dd, 1 H,  $J = 2.4 \text{ Hz}, 8.2 \text{ Hz}$ ), 5.08 (s, 2 H), 3.88 (t, 2 H,  $J = 6.0 \text{ Hz}$ ), 2.80 - 2.63 (m, 2 H), 2.59 - 2.45 (m, 4 H), 2.10 (s, 3 H), 1.41 - 1.30 (m, 4 H), 0.79 (t, 6 H,  $J = 7.3 \text{ Hz}$ ); IR 3400, 2900, 1250; MS FAB m/z 473 [M+H+]; CHN calcd for  $\text{C}_{30}\text{H}_{36}\text{N}_2\text{O}_3 + .20 \text{ H}_2\text{O}$ .

30 Example No. 103 1-[4-(2-Dibutylamino-ethoxy)-benzyl]-2-(4-hydroxy-phenyl)-3-methyl-1H-indol-5-ol

Foam;  $^1\text{H}$  NMR (DMSO) 9.63 (s, 1H), 8.66 (s, 1 H), 7.15 (d, 2 H,  $J = 8.6 \text{ Hz}$ ), 7.05 (d, 1 H,  $J = 8.8 \text{ Hz}$ ), 6.83 (d, 2 H,  $J = 8.6 \text{ Hz}$ ), 6.79 (d, 1 H,  $J = 4.2 \text{ Hz}$ ), 6.78 - 6.71 (m, 4 H), 6.55

(dd, 1 H, J = 8.6 Hz, J = 2.4 Hz), 5.10 (s, 2 H), 3.88 (t, 2 H, J = 5.5 Hz), 2.68-2.62 (m, 2H), 2.42-2.34 (m, 4 H), 2.08 (s, 3 H), 1.38 - 1.19 (m, 8H), 0.82 (t, 6 H, J = 7.2 Hz); IR (KBr) 3400, 1450 cm<sup>-1</sup>; MS eI m/z 501 (M+).

5 Example No. 104 1-[4-(2-Diisopropylamino-ethoxy)-benzyl]-2-(4-hydroxy-phenyl)-3-methyl-1H-indol-5-ol

Mp = 96 - 102°C; <sup>1</sup>H NMR (DMSO) 9.64 (s, 1 H), 8.67 (s, 1 H), 7.14 (d, 2 H, J = 8.6 Hz), 7.04 (d, 1 H, J = 8.6 Hz), 6.83 (d, 2 H, J = 8.6 Hz), 6.79 (d, 1 H, J = 2.4 Hz), 6.77 - 6.69 (m, 4 H), 6.56 (dd, 1 H, J = 8.6 Hz, 2.2 Hz), 5.08 (s, 2 H), 3.75 (t, 2 H, J = 7.0 Hz), 3.01 - 2.92 (m, 2 H), 2.67 (t, 2 H, J = 7.0 Hz), 2.09 (s, 3 H), 0.93 (d, 12 H, 6.6 Hz); IR (KBr) 3400 br, 2940, 1620 cm<sup>-1</sup>; MS FAB m/z 473 (M+H+); CHN calcd for C<sub>30</sub>H<sub>36</sub>N<sub>2</sub>O<sub>3</sub> + 0.5 H<sub>2</sub>O.

15 Example No. 105 1-[4-[2-(Butyl-methyl-amino)-ethoxyl-benzyl]-2-(4-hydroxy-phenyl)-3-methyl-1H-indol-5-ol

Mp = 102-107°C; <sup>1</sup>H NMR (DMSO) 9.60 (s, 1H), 8.67 (s, 1H), 7.14 (d, 2 H, J = 8.4 Hz), 7.04 (d, 1 H, J = 8.6 Hz), 6.82 (d, 2 H, J = 8.8 Hz), 6.78 (d, 1 H, J = 2.3 Hz), 6.73 (s, 4 H), 6.55 (dd, 1 H, J = 8.8 Hz, J = 2.4 Hz), 5.08 (s, 2 H), 3.92 (t, 2H, J = 6.0 Hz), 2.64-2.59 (m, 2 H), 2.38-2.29(m, 2 H), 2.20 (br s, 3 H), 2.08 (s, 3 H), 1.40-1.31 (m, 2 H), 1.25-1.19 (m, 2 H), 0.83 (t, 3 H, 7.2 Hz); IR (KBr) 3420, 1460, 1230 cm<sup>-1</sup>; MS eI m/z 638 (M+).

25 Example No. 108 2-(4-Hydroxy-phenyl)-3-methyl-1-[4-[2-(2-methyl-piperidin-1-yl)-ethoxyl-benzyl]-1H-indol-5-ol

Mp = 121 - 123°C; <sup>1</sup>H NMR (DMSO) 9.65 (s, 1 H), 8.68 (s, 1 H), 7.14 (d, 2 H, J = 8.6 Hz), 7.04 (d, 1 H, J = 8.8 Hz), 6.84 (d, 2 H, J = 8.6 Hz), 6.79 (d, 1 H, J = 2.0 Hz), 6.74 (s, 4 H), 6.56 (dd, 1 H, J = 8.8 Hz, 2.4 Hz), 5.09 (s, 2 H), 3.97 - 3.86 (m, 2 H), 2.95 - 2.73 (m, 2 H), 2.62 - 2.53 (m, 1 H), 2.36 - 2.14 (m, 2 H), 2.09 (s, 3 H), 1.61 - 1.30 (m, 4 H), 1.28 - 1.09 (m, 2 H), 0.98 (d, 3 H, J = 5.1 Hz); IR (KBr) 3400, 2920, 2850, 1610 cm<sup>-1</sup>; CHN calcd for C<sub>30</sub>H<sub>34</sub>N<sub>2</sub>O<sub>3</sub> + 0.25 H<sub>2</sub>O.

Example No. 109 2-(4-Hydroxy-phenyl)-3-methyl-1-[4-[2-(3-methyl-piperidin-1-yl)-ethoxyl-benzyl]-1H-indol-5-ol

Mp = 121 - 123°C; <sup>1</sup>H NMR (DMSO) 9.64 (s, 1 H), 8.67 (s, 1 H), 7.14 (dd, 2 H, J = 8.6 Hz, 1.4 Hz), 7.04 (dd, 1 H, J = 8.6 Hz, 1.2 Hz), 6.84 (dd, 2 H, J = 8.6 Hz, 1.7 Hz), 6.79 (s, 1 H), 6.79 (s, 4 H), 6.56 (d, 1 H, J = 8.6 Hz), 5.08 (s, 2 H), 3.94 (t, 2 H, J = 5.0 Hz), 2.86 - 2.71 (m, 2 H), 2.63 - 2.50 (m, 2 H), 2.48 (s, 3 H), 1.92 - 1.79 (m, 2 H), 1.63 - 1.35 (m, 5 H), 0.79 (d, 3 H, J = 5.2 Hz); IR (KBr) 3400, 2910, 1625 cm<sup>-1</sup>; CHN calcd for C<sub>30</sub>H<sub>34</sub>N<sub>2</sub>O<sub>3</sub> + 0.25 H<sub>2</sub>O.

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Example No. 110 2-(4-Hydroxy-phenyl)-3-methyl-1-[4-[2-(4-methyl-piperidin-1-yl)-ethoxyl-benzyl]-1H-indol-5-ol (HCl)

Mp = 154 - 162°C; <sup>1</sup>H NMR (DMSO) 10.00 (brs, 1 H), 9.71 (s, 1 H), 8.71 (s, 1 H), 7.15 (d, 2 H, J = 8.6 Hz), 7.05 (d, 1 H, J = 8.6 Hz), 6.85 (d, 2 H, J = 8.6 Hz), 6.83 - 6.77 (m, 4 H), 6.57 (dd, 1 H, J = 8.6 Hz, 2.2 Hz), 5.11 (s, 2 H), 4.27 (t, 2 H, J = 4.8 Hz), 3.51 - 3.35 (m, 4 H), 3.01 - 2.87 (m, 2 H), 2.09 (s, 3 H), 1.74 (d, 2 H, J = 13.4 Hz), 1.61 - 1.37 (m, 4 H), 0.88 (d, 3 H, J = 6.4 Hz); IR (KBr) 3410, 2910, 1620 cm<sup>-1</sup>; MS ei m/z 470 (M+H<sup>+</sup>); CHN calcd for C<sub>30</sub>H<sub>34</sub>N<sub>2</sub>O<sub>3</sub> + HCl + 2 H<sub>2</sub>O.

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Example No. 111 1-[4-[2-(3,3-Dimethyl-piperidin-1-yl)-ethoxyl-benzyl]-2-(4-hydroxy-phenyl)-3-methyl-1H-indol-5-ol

Mp = 100°C; <sup>1</sup>H NMR (DMSO) 9.65 (s, 1 H), 8.67 (s, 1 H), 7.15 (d, 2 H, J = 8.6 Hz), 7.05 (d, 1 H, J = 8.8 Hz), 6.84 (d, 2 H, J = 8.6 Hz), 6.79 (d, 1 H, J = 2.4 Hz), 6.74 (s, 4 H), 6.56 (dd, 1 H, J = 8.8, 2.4 Hz), 5.09 (s, 2 H), 3.93 (t, 2 H, J = 5.7 Hz), 2.60-2.50 (m, 2 H), 2.37-2.25 (m, 2 H), 2.09 (s, 3 H), 2.10-1.99 (m, 2 H), 1.46 (t, 2 H, J = 5.9 Hz), 1.13 (t, 2 H, J = 6.4 Hz), 0.86 (s, 6 H); MS ei m/z 484.

30 Example No. 112 1-[4-[2-((cis)-2,6-Dimethyl-piperidin-1-yl)-ethoxyl-benzyl]-2-(4-hydroxy-phenyl)-3-methyl-1H-indol-5-ol

Mp = 114 - 121°C; <sup>1</sup>H NMR (DMSO) 9.62 (s, 1 H), 8.64 (s, 1 H), 7.11 (d, 2 H, J = 8.6 Hz), 7.01 (d, 1 H, J = 8.6 Hz), 6.81 (d, 2 H, J = 8.8 Hz), 6.76

(d, 1 H,  $J = 2.2$  Hz), 6.72 - 6.66 (m, 4 H), 6.53 (dd, 1 H,  $J = 8.6$  Hz, 2.2 Hz), 5.06 (s, 2 H), 3.86 - 3.72 (m, 2 H), 2.86 - 2.76 (m, 2 H), 2.43 - 2.35 (m, 2 H), 2.06 (s, 3 H), 1.78 - 1.59 (m, 3 H), 1.29 - 1.17 (m, 1 H), 1.12 - 0.92 (m, 8 H); IR (KBr) 3400 br, 2920, 1630  $\text{cm}^{-1}$ ; MS FAB m/z 485 ( $M+H^+$ ); CHN calcd for  $\text{C}_{31}\text{H}_{36}\text{N}_2\text{O}_3$  + 5 0.1 acetone + 0.75  $\text{H}_2\text{O}$ .

Example No. 113 2-(4-Hydroxy-phenyl)-1-[4-[2-(4-hydroxy-piperidin-1-yl)-ethoxyl-benzyl]-3-methyl-1H-indol-5-ol  
Mp = 80 - 90°C;  $^1\text{H}$  NMR (DMSO) 9.66 (s, 1 H), 8.68 (s, 1 H), 7.15 (d, 2 H,  $J = 7.6$  Hz), 7.04 (d, 1 H,  $J = 8.8$  Hz), 6.84 (dd, 2 H,  $J = 2.0$  Hz, 6.6 Hz), 6.78 (d, 1 H, 2.2 Hz), 6.73 (s, 4 H), 6.55 (dd, 1 H,  $J = 2.2$  Hz, 8.6 Hz), 5.09 (s, 2 H), 4.50 (d, 1 H,  $J = 4.2$  Hz), 3.92 (t, 2 H,  $J = 5.8$  Hz), 3.40 (m, 2 H), 2.72 (m, 2 H), 2.60 (m, 2 H), 2.10 (s, 3 H), 2.15-2.05 (m, 1 H), 1.75-1.63 (m, 2 H), 1.42 - 1.28 (m, 2 H); IR (KBr) 3400, 2900, 1250  $\text{cm}^{-1}$ ; MS eI m/z 472 ( $M^+$ ); CHN calcd for  $\text{C}_{29}\text{H}_{32}\text{N}_2\text{O}_4$  + .11  $\text{CH}_2\text{Cl}_2$ .

Example No. 114 (1S,4R)-1-[4-[2-(2-Aza-bicyclo [2.2.1] hept-2-yl)-ethoxyl-benzyl]-2-(4-hydroxy-phenyl)-3-methyl-1H-indol-5-ol  
Mp = 125 - 130°C;  $^1\text{H}$  NMR (DMSO) 9.65 (s, 1 H), 8.67 (s, 1 H), 7.13 (d, 2 H,  $J = 8.6$  Hz), 7.04 (d, 1 H,  $J = 8.5$  Hz), 6.83 (dd, 2 H,  $J = 2.0$  Hz, 6.6 Hz), 6.78 (d, 1 H,  $J = 2.2$  Hz), 6.73 (s, 4 H), 6.55 (dd, 1 H,  $J = 2.2$  Hz, 8.6 Hz), 5.08 (s, 2 H), 3.95 - 3.8 (m, 2 H), 2.90 - 2.70 (3 H), 2.30 - 2.20 (m, 2 H), 2.10 (s, 3 H), 1.70 - 1.60 (m, 1 H), 1.60 - 1.30 (m, 4 H), 1.25 - 1.15 (m, 2 H); IR (KBr) 3400, 2950, 1500; MS (+) FAB m/z 469 [ $M+H]^+$ ; CHN calcd for  $\text{C}_{30}\text{H}_{32}\text{N}_2\text{O}_3$  + .34 EtOAc.

Example No. 115 2-(4-Hydroxy-phenyl)-3-methyl-1-[4-[2-(1,3,3-trimethyl-6-aza-bicyclo[3.2.1]oct-6-yl)-ethoxyl-benzyl]-1H-indol-5-ol  
Mp = 98 - 100°C;  $^1\text{H}$  NMR (DMSO) 9.64 (s, 1 H), 8.67 (s, 1 H), 7.14 (d, 2 H,  $J = 8.6$  Hz), 7.05 (d, 1 H,  $J = 8.6$  Hz), 6.84 (d, 2 H,  $J = 8.6$  Hz), 6.79 (d, 1 H,  $J = 2.4$  Hz), 6.75 - 6.69 (m, 4 H), 6.56 (dd, 1 H,  $J = 8.6$  Hz, 2.4 Hz), 5.08 (s, 2 H), 3.83 (t, 2 H,  $J = 5.9$  Hz), 3.12 - 3.07 (m, 1 H), 2.94 - 2.87 (m, 1 H), 2.85 (d, 1 H,  $J = 9.2$  Hz), 2.78 - 2.70 (m, 1 H), 2.17 (d, 1 H,  $J = 9.2$  Hz), 2.09 (s, 3 H), 1.55 - 1.42 (m, 2 H), 1.29 (q, 2 H,  $J = 13.6$  Hz), 1.14 (s, 3 H), 1.11 - 1.02 (m, 2 H),

0.96 (s, 3 H), 0.82 (s, 3 H); IR (KBr) 3400 br, 2940, 2900, 1630  $\text{cm}^{-1}$ ; MS ESI m/z 525 (M+H+); CHN calcd for  $\text{C}_{34}\text{H}_{40}\text{N}_2\text{O}_3 + 0.5 \text{H}_2\text{O}$ .

5 Example No. 116 2-(4-Fluoro-phenyl)-3-methyl-1-[4-(2-piperidine-1-yl-ethoxy)-benzyl]-1H-indol-5-ol (HCl)

Mp = 201 - 203°C;  $^1\text{H}$  NMR (DMSO) 10.22 (s, 1 H), 8.78 (s, 1 H), 7.45 - 7.35 (m, 2 H), 7.34 - 7.25 (m, 2 H), 7.11 (d, 1 H,  $J$  = 8.6 Hz), 6.90 - 6.70 (m, 5 H), 6.61 (dd, 1 H,  $J$  = 2.4 Hz, 8.8 Hz), 5.15 (s, 2 H), 4.27 (t, 2 H, 4.8 Hz), 3.50 - 3.34 (m, 4 H), 3.0 - 2.85 (m, 2 H), 2.10 (s, 3 H), 1.80 (m, 5 H), 1.40 - 1.25 (m, 1 H);  
10 MS eI m/z 458 (M+); CHN calcd for  $\text{C}_{29}\text{H}_{31}\text{FN}_2\text{O}_2 + 1 \text{HCl}$ .

15 Example No. 117 1-[4-(2-Azepan-1-yl-ethoxy)-benzyl]-2-(4-fluoro-phenyl)-3-methyl-1H-indol-5-ol

Mp = 181 - 184°C;  $^1\text{H}$  NMR (DMSO) 10.68 (s, 1 H), 8.80 (s, 1 H), 7.50 - 7.36 (m, 2 H), 7.34 - 7.26 (m, 2 H), 7.12 (d, 1 H,  $J$  = 8.8 Hz), 6.86 - 6.73 (m, 5 H), 6.63 (dd, 1 H,  $J$  = 2.2 Hz, 8.5 Hz), 5.13 (s, 2H), 4.29 (t, 2 H,  $J$  = 5.2 Hz), 3.50 - 3.30 (m, 4 H), 3.20 - 3.08 (m, 2 H), 2.11 (s, 3 H), 1.90 - 1.70 (m, 4 H), 1.68 - 1.45 (m, 4 H); IR (KBr) 3500, 3100, 2910, 1450, 1250  $\text{cm}^{-1}$ ; MS eI m/z 472 (M+); CHN calcd for  $\text{C}_{30}\text{H}_{33}\text{FN}_2\text{O}_2 + 1 \text{HCl}$ .

20 Example No. 118 2-(3-Methoxy-4-hydroxy-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indol-5-ol (HCl)

Mp = 161-163°C;  $^1\text{H}$  NMR (DMSO) 10.12 (brs, 1H), 9.25 (s, 1 H), 8.71 (s, 1H), 7.05 (d, 1H,  $J$  = 8.5Hz), 6.85 - 6.79 (m, 8 H), 6.57 (dd, 1H,  $J$  = 8.5Hz,  $J$  = 2.2Hz), 5.13 (s, 2H), 4.27 (t, 2H,  $J$  = 5.0Hz), 3.64 (s, 3H), 3.44 - 3.37 (m, 4 H), 2.93 - 2.85 (m, 2H), 2.11 (s, 3H), 1.80 - 1.60 (m, 5 H), 1.40 - 1.25 (m, 1H); MS eI m/z 486 (M+); CHN calc for  $\text{C}_{30}\text{H}_{34}\text{N}_2\text{O}_4 + 1\text{HCl} + 1\text{H}_2\text{O}$ ; IR (KBr) 3190, 1470, 1230  $\text{cm}^{-1}$ .

30 Example No. 119 2-Benzof[1,3]dioxol-5-yl-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indol-5-ol (HCl)

Mp = 122-125°C;  $^1\text{H}$  NMR (DMSO) 9.80 (brs, 1 H), 8.73 (s, 1 H), 7.07

(d, 1 H, J = 8.7 Hz), 7.02 (d, 1 H, J = 8.0 Hz), 6.89 (d, 1 H, J = 1.7 Hz), 6.80 - 6.75 (m, 6 H), 6.58 (dd, 1 H, J = 6.4 Hz, J = 2.2 Hz), 6.06 (s, 2 H), 5.13 (s, 2 H), 4.30 - 4.19 (m, 2 H), 3.51 - 3.30 (m, 4 H), 2.99-2.85 (m, 2 H), 2.10 (s, 3 H), 1.81-1.59 (m, 5 H), 1.41-1.26 (m, 1 H); MS eI m/z 484(M+); CHN calc for  
5  $C_{30}H_{32}N_2O_4$  + HCl + .26 H<sub>2</sub>O.

Example No. 120 2-(4-Isopropoxy-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indol-5-ol (HCl)

10 Mp = 120 - 125°C; <sup>1</sup>H NMR (DMSO) 10.18 (s, 1 H), 8.73 (s, 1 H), 7.25 (d, 2 H, J = 8.6 Hz), 7.04 (d, 1 H, J = 8.8 Hz), 6.99 (d, 2 H, J = 8.8 Hz), 6.82 - 6.80 (m, 5 H), 6.59 (dd, 1 H, J = 2.2 Hz, 8.6 Hz), 5.12 (s, 2 H), 4.67 - 4.61 (m, 1 H), 4.27 (t, 2 H, J = 4.8 Hz), 3.50 - 3.35 (m, 4 H), 3.0 - 2.85 (m, 2 H), 2.10 (s, 3 H), 1.80 - 1.60 (m, 5 H), 1.40 - 1.25 (m, 7 H); IR (KBr) 3400, 3000, 1500, 1250; MS eI m/z 498 (M+); CHN calcd for  $C_{32}H_{38}N_2O_3$  + 1.0 HCl + .70 H<sub>2</sub>O.

15

Example No. 121 1-[4-(2-Azepan-1-yl-ethoxy)-benzyl]-2-(4-isopropoxy-phenyl)-3-methyl-1H-indol-5-ol (HCl)

20 Mp = 120 - 125°C; <sup>1</sup>H NMR (DMSO) 10.36 (s, 1 H), 8.73 (s, 1 H), 7.26 - 7.23 (m, 2 H), 7.05 (d, 1 H, J = 8.8 Hz), 7.01 - 6.98 (m, 2 H), 6.85 - 6.75 (m, 5 H), 6.57 (dd, 1 H, J = 2.2 Hz, 8.6 Hz), 5.12 (s, 2 H), 4.67 - 4.61 (m, 1 H), 4.27 (t, 2 H, J = 4.8 Hz), 3.50 - 3.30 (m, 4 H), 3.20 - 3.10 (m, 2 H), 2.10 (s, 3 H), 1.85 - 1.75 (m, 4 H), 1.65 - 1.50 (m, 4 H), 1.27 (d, 6 H, J = 6.1 Hz); IR (KBr) 3400, 1500, 1250; MS eI m/z 512 (M+); Calcd for  $C_{33}H_{40}N_2O_3$  + 1.0 HCl + .5 H<sub>2</sub>O.

25 Example No. 122 2-(4-Cyclopropylmethoxy-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indol-5-ol

30 Mp = 121 - 135°C; <sup>1</sup>H NMR (DMSO) 9.80 (br s, 1 H), 8.72 (s, 1 H), 7.24 (d, 2 H, J = 8.8 Hz), 7.05 (d, 1 H, J = 8.8 Hz), 6.98 (d, 2 H, J = 8.8 Hz), 6.83 - 6.78 (m, 5 H), 6.57 (dd, 1 H, J = 8.8 Hz, 2.4 Hz), 5.13 (s, 2 H), 4.86 - 4.82 (m, 1 H), 4.25 (t, 2 H, J = 4.8 Hz), 3.50 - 3.38 (m, 4 H), 2.92 (q, 2 H, J = 8.8 Hz), 2.11 (s, 3 H), 1.98 - 1.85 (m, 2 H), 1.81 - 1.56 (m, 11 H), 1.41 - 1.29 (m, 1 H); IR (KBr) 3400, 2920, 1620 cm<sup>-1</sup>; MS eI m/z 524 (M+); CHN calcd for  $C_{34}H_{40}N_2O_3$  + 0.5 H<sub>2</sub>O.

Example No. 123 3-Methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-2-(4-trifluoromethyl-phenyl)-1H-indol-5-ol

5 Mp = 174°C;  $^1\text{H}$  NMR (DMSO) 8.8 (s, 1 H), 7.82 (d, 2 H, J = 8.1 Hz), 7.59 (d, 2 H, J = 7.9 Hz), 7.17 (d, 1 H, J = 8.6 Hz), 6.86 (d, 1 H, J = 2.4 Hz), 6.75 - 6.68 (m, 4 H), 6.65 (dd, 1 H, J = 8.8 Hz, 2.4 Hz), 5.16 (s, 2 H), 3.93 (t, 2 H, J = 5.7 Hz), 2.62 - 2.56 (m, 2 H), 2.42 - 2.32 (m, 4 H), 2.15 (s, 3 H), 1.48 - 1.40 (m, 4 H), 1.39 - 1.29 (m, 2 H); IR (KBr) 3410, 2910, 2850, 1620 cm<sup>-1</sup>; MS eI m/z 508 (M $^+$ ); CHN calcd for  $\text{C}_{30}\text{H}_{31}\text{F}_3\text{N}_2\text{O}_2$  + 0.25  $\text{H}_2\text{O}$ .

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Example No. 124 3-Methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-2-p-tolyl-1H-indol-5-ol

15 Mp = 162 - 164°C;  $^1\text{H}$  NMR (DMSO) 8.70 (s, 1 H), 7.28 - 7.24 (m, 4 H), 7.07 (d, 1 H, J = 8.4 Hz), 6.81 (d, 1 H, J = 2.2 Hz), 6.73 (s, 4 H), 6.58 (dd, 1 H, J = 2.4 Hz, 8.8 Hz), 5.11 (s, 2 H), 3.92 (t, 2 H, J = 5.9 Hz), 2.55 (t, 2 H, J = 5.9 Hz), 2.45 - 2.30 (m, 7 H), 2.10 (s, 3 H), 1.50 - 1.40 (m, 4 H), 1.48 - 1.35 (m, 2 H); IR (KBr) 3400, 2900, 1200; MS eI m/z 454 (M $^+$ ); CHN calcd for  $\text{C}_{30}\text{H}_{34}\text{N}_2\text{O}_2$ .

20 Example No. 125 2-(4-Chloro-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indol-5-ol (HCl)

Mp = 161-164°C;  $^1\text{H}$  NMR (DMSO) 10.12 (brs, 1H), 8.80 (s, 1H), 7.53 (d, 2H, J = 8.3 Hz), 7.36 (d, 2H, J = 8.8 Hz), 7.12 (d, 1 H, J = 8.8 Hz), 6.85 - 6.75 (m, 5 H), 6.63 (dd, 1H, J = 8.8 Hz, J = 2.4 Hz), 5.14 (s, 2H), 4.29-4.22 (m, 2H), 3.45-3.36 (m, 4 H), 2.97 - 2.84 (m, 2H), 2.11 (s, 3H), 1.83 - 1.61 (m, 5H), 1.37 - 1.25 (m, 1H); MS eI m/z 475 (M $^+$ ); CHN calc for  $\text{C}_{29}\text{H}_{31}\text{ClN}_2\text{O}_2$  + HCl + .25  $\text{H}_2\text{O}$ .

Example No. 126 2-(2,4-Dimethoxy-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indol-5-ol

30 Mp = 85 - 92°C;  $^1\text{H}$  NMR (DMSO) 8.62 (s, 1 H), 7.10 (d, 1 H, J = 8.4 Hz), 7.01 (d, 1 H, J = 8.6 Hz), 6.80 - 6.70 (m, 5 H), 6.69 (d, 1 H, 2.2 Hz), 6.59 (dd, 1 H, J = 2.4 Hz, 8.5 Hz), 6.52 (dd, 1 H, J = 2.4 Hz, 8.8 Hz), 5.02

(d, 1 H, J = 6.5 Hz), 4.83 (d, 1 H, J = 6.3 Hz), 4.0 - 3.90 (m, 2 H), 3.80 (s, 3 H), 3.67 (s, 3 H), 2.65 - 2.50 (m, 2 H), 2.45 - 2.30 (m, 4 H), 2.0 (s, 3 H), 1.55 - 1.40 (m, 4 H), 1.39 - 1.30 (m, 2 H); IR (KBr) 3400, 2900, 1520, 1250; MS eI m/z 500 (M+); CHN calcd for  $C_{31}H_{36}N_2O_4$  + .05  $CH_2Cl_2$ .

5

**Example No. 127 2-(3-Hydroxy-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indol-5-ol**

Mp = 115 - 118°C;  $^1H$  NMR (DMSO) 9.57 (s, 1 H), 8.71 (s, 1 H), 7.27 - 7.23 (t, 1 H, J = 8.1 Hz), 7.06 - 7.04 (d, 1 H, J = 8.8 Hz), 6.81 - 6.74 (m, 8 H), 6.59 - 6.56 (dd, 1 H, J = 2.3 Hz, J = 6.3 Hz), 5.12 (s, 2 H), 3.94 - 3.91 (t, 2 H, J = 5.9 Hz), 2.57 - 2.54 (t, 2 H, J = 5.8 Hz), 2.36 (s, 4 H), 2.11 (s, 3 H), 1.45 - 1.41 (m, 4 H), 1.34 - 1.33 (m, 2 H); IR (KBr) 3400, 2900  $cm^{-1}$ ; MS eI m/z 456 (M+); CHN calcd for  $C_{29}H_{32}N_2O_3$  + 1.0  $H_2O$ .

15

**Example No. 128 1-[4-(2-Azepan-1-yl-ethoxy)-benzyl]-2-(3-hydroxy-phenyl)-3-methyl-1H-indole-5-ol**

Mp = 94 - 97°C;  $^1H$  NMR (DMSO) 9.58 (s, 1 H), 8.71 (s, 1 H), 7.27 - 7.23 (t, 1 H, J = 7.9 Hz), 7.07 - 7.04 (d, 1 H, J = 8.7 Hz), 6.81 - 6.74 (m, 8 H), 6.59 - 6.56 (dd, 1 H, J = 2.4 Hz, J = 6.3 Hz), 5.12 (s, 2 H), 3.9 (m, 2 H), 2.80 (s, 2 H), 2.65 (s, 4 H), 2.11 (s, 3 H), 1.54 - 1.50 (m, 8 H); IR 3400, 2900  $cm^{-1}$ ; MS eI m/z 470 (M+); CHN calcd for  $C_{30}H_{34}N_2O_3$  + 0.75  $H_2O$  + 0.23 Ethyl Acetate.

**Example No. 129 2-(3-Fluoro-4-hydroxy-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indol-5-ol**

25 Mp = 117-119°C;  $^1H$  NMR (DMSO) 10.1 (s, 1 H), 8.71 (s, 1 H), 7.10 - 6.95 (m, 4 H), 6.80 (d, 1 H, J = 2.2 Hz), 6.74 (s, 4 H), 6.59 (dd, 1 H, J = 2.2 Hz, 8.5 Hz), 5.1 (s, 2 H), 3.93 (t, 2 H, J = 5.9 Hz), 2.56 (t, 2 H, J = 5.8 Hz), 2.44 - 2.30 (m, 4 H), 2.10 (s, 3 H), 1.45 - 1.40 (m, 4 H), 1.36 - 1.32 (m, 2 H); MS eI m/Z 475 (M+); CHN calcd for  $C_{29}H_{31}FN_2O_3$ .

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Example No. 130 2-(3-Fluoro-4-hydroxy-phenyl)-3-methyl-1-[4-(azepan-1-yl-ethoxy)-benzyl]-1H-indol-5-ol

10 Mp = 88 - 91°C;  $^1\text{H}$  NMR (DMSO) 10.10 (s, 1H), 8.71 (s, 1H), 7.12 - 6.94 (m, 4 H), 6.80 (d, 1 H,  $J$  = 2.2 Hz), 6.74 (s, 4 H), 6.58 (dd, 1 H,  $J$  = 2.2 Hz, 8.5 Hz), 5.10 (s, 2 H), 3.91 (t, 2 H,  $J$  = 5.9 Hz), 2.76 (t, 2 H,  $J$  = 5.9), 2.62 - 2.60 (m, 4H), 2.10 (s, 3H), 1.70 - 1.40 (m, 8 H); MS eI m/Z 488 (M+); CHN calcd for  $\text{C}_{30}\text{H}_{33}\text{FN}_2\text{O}_3$ .

10 Example No. 131 2-(3-Methoxy-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole-5-ol

15 Mp = 120 - 123°C;  $^1\text{H}$  NMR (DMSO) 8.76 (s, 1 H), 7.42 - 7.46 (t, 1 H,  $J$  = 7.9 Hz), 7.12 - 7.09 (d, 1 H,  $J$  = 8.7 Hz), 6.99 - 6.92 (m, 2 H), 6.86 - 6.83 (m, 2 H), 6.76 (s, 4 H), 6.63 - 6.60 (dd, 1 H,  $J$  = 2.1 Hz,  $J$  = 6.5 Hz), 5.14 (s, 2 H), 3.96 - 3.92 (t, 2 H,  $J$  = 5.9 Hz), 3.70 (s, 3 H), 2.59 - 2.55 (t, 2 H,  $J$  = 5.9 Hz), 2.37 (s, 4 H), 2.14 (s, 3 H), 1.49 - 1.44 (m, 4 H), 1.35 - 1.34 (m, 2 H); IR 3400, 2950, 1600  $\text{cm}^{-1}$ ; MS eI m/z 471 (M+); CHN calcd for  $\text{C}_{30}\text{H}_{34}\text{N}_2\text{O}_3$ .

20 Example No. 132 3-Methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-2-(4-trifluoromethoxy-phenyl)-1H-indole-5-ol

25 Mp = 122 - 125°C;  $^1\text{H}$  NMR (DMSO) 8.80 (s, 1 H), 7.51 - 7.45 (m, 4 H), 7.17 - 7.14 (d, 1 H,  $J$  = 8.7 Hz), 6.85 - 6.84 (d, 1 H,  $J$  = 2.0 Hz), 6.75 - 6.69 (m, 4 H), 6.66 - 6.62 (m, 1 H), 5.14 (s, 2 H), 3.95 - 3.92 (t, 2 H,  $J$  = 5.8 Hz), 2.59 - 2.55 (t, 2 H,  $J$  = 5.6 Hz), 2.49 - 2.38 (m, 4 H), 2.13 (s, 3 H), 1.47 - 1.44 (m, 4 H), 1.36 - 1.34 (d, 2 H,  $J$  = 4.8 Hz); IR 3400, 2900, 1600  $\text{cm}^{-1}$ ; MS eI m/z 525 (M+); CHN calcd for  $\text{C}_{30}\text{H}_{31}\text{F}_3\text{N}_2\text{O}_3 + 0.25 \text{H}_2\text{O}$ .

Synthetic procedures and physical data for compounds substituted with chloro, ethyl or cyano groups at the 3-position of the indole

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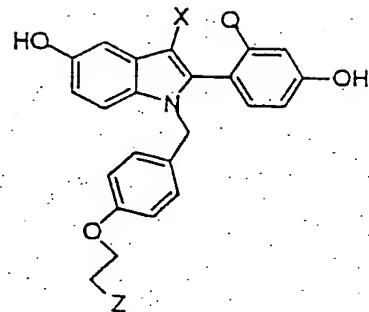


Table 8

Example No.	X	Q	Z
No. 133	Cl	H	N
No. 134	Cl	H	N
No. 135	Cl	H	N
No. 136	Cl	CH <sub>3</sub>	N
No. 137	Et	H	N
No. 138	CN	H	N
No. 139	CN	H	N

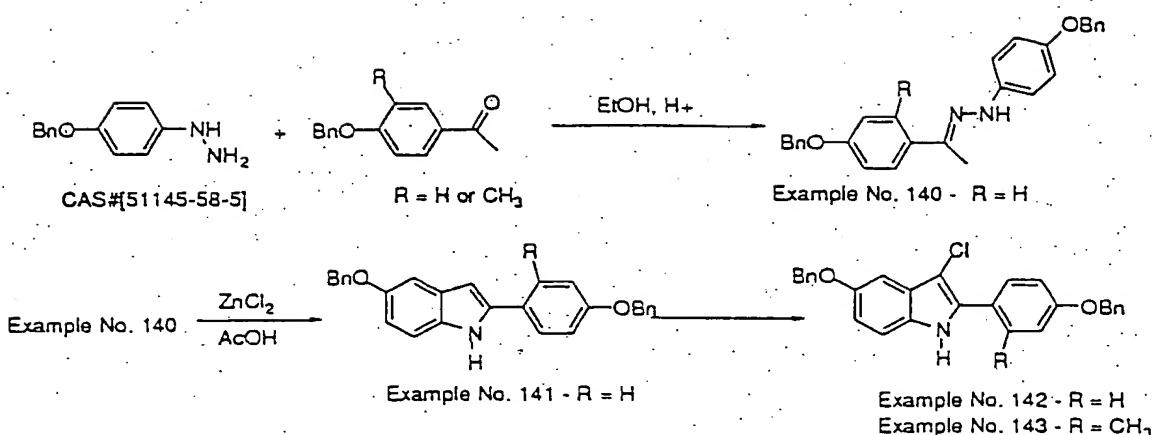
5

Synthesis of 3-chloro analogues No. 133- No. 136

Scheme 14

10 Synthesis of 3-chloroindole

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**Example No. 140. Formation of hydrazone**

5 4-Benzylxyphenylhydrazine CAS No. [51145-58-5] (50.0 g, 233.4 mmol) was mixed with 4-benzylxyacetophenone CAS No. [54696-05-8] (63.0 g, 280.0 mmol) in pure ethanol (800 mL). A catalytic amount of acetic acid (5 drops) was added. The reaction was heated to reflux for 2.5 hrs. During the course of refluxing, the condensed product solidified out of the hot solution. The reaction was cooled down to rt. The desired product was collected by vacuum filtration as a light yellow solid (85 g, 86%). Mp = 165-174°C; <sup>1</sup>H NMR (DMSO) 8.91 (s, 1 H), 7.68 (d, 2 H, J = 8.8 Hz), 7.48 - 7.32 (m, 10 H), 7.12 (d, 2 H, J = 9 Hz), 7.00 (d, 2 H, J = 8.8 Hz), 6.88 (d, 2 H, J = 9.0 Hz), 5.11 (s, 2 H), 5.01 (s, 2 H), 2.17 (s, 3 H); MS eI m/z 422 (M<sup>+</sup>).

15 Example No. 141. Formation of indole from hydrazone: 5-Benzylxy-2-(4-benzylxy-phenyl)-1H-indole

20 A flask was charged with N-(4-Benzylxy-phenyl)-N'-(1-(4-benzylxy-phenyl)-ethylidene)-hydrazine (No. 140) (10.0 g, 23.7 mmol), ZnCl<sub>2</sub> (8.06 g, 59.17 mmol), acetic acid (70 mL). The reaction flask was heated to 105 °C for no more than 20 min. During the heating period, the reaction was monitored carefully by TLC for the disappearance of the starting material. The progress of the reaction could be shown as the product solidified out of the solution while heating. The reaction was then cooled to rt and more product crashed out was observed. The reaction content was poured into a

separatory funnel containing ether (100 mL) and H<sub>2</sub>O (200 mL), which was shaken vigorously. The insoluble residue as the desired product stayed in the ether layer which was collected by vacuum filtration. The product was further purified by trituration in ether to give a light gray solid (4.4 g, 46%).

5 Mp = 202 - 204°C; <sup>1</sup>H NMR (DMSO) 11.24 (s, 1 H), 7.73 (d, 2 H, J = 8.8 Hz), 7.48 - 7.41 (m, 4 H), 7.45 - 7.27 (m, 6 H), 7.25 (d, 1 H, J = 8.6 Hz), 7.12 - 7.04 (m, 3 H), 6.77 (dd, 1 H, J = 2.4 Hz, 8.6 Hz), 6.65 (d, 1 H, J = 1.5 Hz), 5.14 (s, 2 H), 5.08 (s, 2 H); IR 3420, 3000, 1625 cm<sup>-1</sup>; MS eI m/z 405 (M+); CHN calcd for C<sub>28</sub>H<sub>23</sub>NO<sub>2</sub> + 0.40 H<sub>2</sub>O.

10

Example No. 142 Chlorination of indole to render 5-Benzylxy-3-chloro-2-(4-benzylxy-phenyl)-1H-indole

15 A flask was charged with 5-Benzylxy-2-(4-benzylxy-phenyl)-1H-indole No. 141 (8.0 g, 20.0 mmole) and CH<sub>2</sub>Cl<sub>2</sub> (50ml). The reaction was cooled to 0°C and n-chlorosuccinimide (2.9g, 22mmole) was added. The reaction was stirred at 0°C for 20min. The reaction was then washed with 10% sodium sulfite solution, dried over MgSO<sub>4</sub>, and concentrated. To the resulting brown solid was added MeOH and the mixture was stirred for 15 min. The solid was filtered to give 6.8g of a tan solid (78%). Mp = 157-160°C; <sup>1</sup>H NMR (DMSO) 11.5 (s, 1 H), 7.80 (d, 2 H, J = 7.0 Hz), 7.42 - 7.28 (m, 11 H), 7.17 (d, 2 H, J = 8.7 Hz), 7.01 (d, 1 H, J = 2.2Hz), 6.88 (dd, 1 H, J = 8.8 Hz, J = 2.4 Hz), 5.17 (s, 2H), 5.13 (s, 2H); MS eI m/z 439 (M+).

25 Example No. 143 5-Benzylxy-3-chloro-2-(2-methyl-4-benzylxy-phenyl)-1H-indole

This indole synthesized analogously to indole No. 142 immediately preceding: Mp = <sup>1</sup>H NMR (DMSO) 11.34 (s, 1 H), 7.48 - 7.44 (m, 4 H), 7.42 - 7.24 (m, 8 H), 7.02 (dd, 2 H, J = 9.3 Hz, J = 2.4 Hz), 6.95 (dd, 1 H, J = 8.4 Hz, J = 2.6Hz), 6.88 (dd, 1 H, J = 8.8Hz, J = 2.4 Hz), 5.16 (s, 2 H), 5.14 (s, 2 H), 2.23 (s, 3 H); MS eI m/z 453 (M+).

Example No. 144 Alkylation of indole to give {4-[5-Benzyl-2-(4-benzyl-phenyl)-3-chloro-indol-1-ylmethyl-phenoxyl]-acetic acid ethyl ester}

5

This procedure was performed analogously to that outlined for the synthesis of 3-methyl indole acetic acid ethyl esters outlined in method 3.

Mp = 90-94°C;  $^1\text{H}$  NMR (DMSO) 7.45 (d, 4H,  $J = 7.3$  Hz), 7.41 - 7.26 (m, 9H), 7.14 (d, 2H,  $J = 8.7$  Hz), 7.04 (d, 1H,  $J = 2.4$  Hz), 6.91

10 (dd, 1H,  $J = 9.0$  Hz,  $J = 2.5$  Hz), 6.80-6.74 (m, 4H), 5.24 (s, 2H), 5.15 (s, 2H), 5.14 (s, 2H), 4.66 (s, 2H), 4.12 (q, 2H,  $J = 7.2$  Hz), 1.16 (t, 3H,  $J = 7.5$  Hz); MS eI m/z 631(M+).

Example No. 145 Reduction of No. 144 to render No. 145 2-[4-[5-

15 Benzyl-2-(4-benzyl-phenyl)-3-chloro-indol-1-ylmethyl-phenoxyl]-ethanol

This reaction was performed analogously to that outlined for the synthesis of 3-methyl indoles outlined in method 4. Compound was not purified or characterized, but used as obtained for the next step.

Example No. 146 Bromination of No. 145 to render Benzyl-2-(4-benzyl-phenyl)-1-[4-(2-bromo-ethoxy)-benzyl]-3-chloro-1H-indole

25 This reaction was performed analogously to that outlined for the synthesis of 3-methyl indoles outlined in method 5. Mp = 155-158°C;  $^1\text{H}$  NMR (DMSO) 7.45 (d, 4H,  $J = 7.8$  Hz), 7.41 - 7.25 (m, 9H), 7.14 (d, 2H,  $J = 8.7$  Hz), 7.04 (d, 1H,  $J = 2.4$  Hz), 6.91 (dd, 1H,  $J = 9.0$  Hz,  $J = 2.5$  Hz), 6.74 (s, 4H), 5.24 (s, 2H), 5.15 (s, 2H), 5.14 (s, 2H), 4.20 (t, 2H,  $J = 5.3$  Hz), 3.74 (t, 2H,  $J = 5.3$  Hz); MS eI m/z 651 (M+).

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Example No. 147 Substitution of No. 146 with piperidine to render 5-Benzyl-2-(4-benzyl-phenyl)-3-chloro-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole

5 This reaction performed analogously to that outlined for the synthesis of 3-methyl indoles outlined in method 6, using piperidine to substitute the bromide.  
Mp 96-98°C;  $^1\text{H}$  NMR (DMSO) 7.45 (d, 4 H,  $J = 7.8$  Hz), 7.40 - 7.30 (m, 9 H), 7.14 (d, 2 H,  $J = 8.7$  Hz), 7.04 (d, 1 H,  $J = 2.4$  Hz), 6.91 (dd, 1 H,  $J = 9.0$  Hz,  $J = 2.5$  Hz), 6.74 (s, 4 H), 5.24 (s, 2 H), 5.15 (s, 2 H), 5.14 (s, 2 H), 3.93 (t, 2 H,  $J = 6.0$  Hz), 2.56 (t, 2 H,  $J = 6.0$  Hz), 2.41-2.32 (m, 4 H), 1.48-1.39 (m, 4 H), 1.38-1.31 (m, 2 H).

10 Example No. 148 5-Benzyl-2-(4-benzyl-phenyl)-3-chloro-1-[4-(2-azepan-1-yl-ethoxy)-benzyl]-1H-indole

15 Reaction performed the same as above except the substituting amine used was hexamethylenearmine.  
Mp = 94-97°C;  $^1\text{H}$  NMR (DMSO) 7.45 (d, 4H,  $J = 7.8$  Hz), 7.42 - 7.30 (m, 9H), 7.14 (d, 2 H,  $J = 8.7$  Hz), 7.04 (d, 1 H,  $J = 2.4$  Hz), 6.91 (dd, 1 H,  $J = 9.0$  Hz,  $J = 2.5$  Hz), 6.74 (s, 4 H), 5.24 (s, 2 H), 5.15 (s, 2 H), 5.14 (s, 2 H), 3.93 (t, 2 H,  $J = 6.0$  Hz), 2.75 (t, 2 H,  $J = 6.0$  Hz), 2.63-2.59 (m, 4 H), 1.58-1.44 (m, 8 H); MS eI m/z 671 (M+).

20 Example No. 149 5-Benzyl-2-(2-methyl-4-benzyl-phenyl)-3-chloro-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole

Reactions to make this compound analogous to those used to make No. 147.  
Oil;  $^1\text{H}$  NMR(DMSO) 7.50 - 7.29 (m, 11 H), 7.17 (d, 1 H,  $J = 8.4$  Hz), 7.05 (d, 1 H,  $J = 2.4$  Hz), 7.02 (d, 1 H,  $J = 2.4$  Hz), 6.93 - 6.85 (m, 2 H), 6.75 - 6.65 (m, 4 H), 5.14 (s, 2 H), 5.13 (s, 2 H), 5.07 (m, 2 H), 3.92 (t, 2 H,  $J = 5.9$  Hz), 2.55 (t, 2 H,  $J = 5.9$  Hz), 2.42 - 2.29 (m, 4 H), 1.94 (s, 3 H), 1.44 - 1.40 (m, 4 H), 1.38 - 1.34 (m, 2 H).

**Example No. 133 3-Chloro-2-(4-hydroxy-phenyl)-1-[4-(2-pyrrolidin-1-yl-ethoxy)-benzyl]-1H-indol-5-ol (HCl)**

Synthesized as described for example No. 134.

5 Mp = 233-235°C; <sup>1</sup>H NMR (DMSO) 10.50 (s, 1 H), 9.88 (s, 1 H), 9.01 (s, 1 H), 7.30 - 7.20 (m, 3 H), 6.90 - 6.80 (m, 7 H), 6.68 (dd, 1 H, J = 2.4, Hz, 8.8 Hz), 5.20 (s, 2 H), 4.22 (t, 2 H, J = 4.8 Hz), 3.47 (t, 2 H, J = 4.8 Hz), 3.10 (bm, 4 H), 1.90 (s, 4 H); IR (KBr) 3400, 1625, 1475, 825 cm<sup>-1</sup>; MS eI m/z 462 (M+); CHN calcd for C<sub>27</sub>H<sub>27</sub>ClN<sub>2</sub>O<sub>3</sub> + 1 HCl + .75 H<sub>2</sub>O.

10 **Example No. 134 Removal of benzyl ethers to render 3-Chloro-2-(4-hydroxy-phenyl)-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indol-5-ol (HCl)**

15 Benzyl ethers were removed analogously to that procedure outlined for 3-methyl indoles outlined in method 7. This compound was then converted to the hydrochloride salt as described previously in method 8; Mp = 207-209°C; <sup>1</sup>H NMR (DMSO) 10.10 (bs, 1 H), 9.86 (s, 1 H), 9.07 (s, 1 H), 7.26 (d, 2 H, J = 8.6 Hz), 7.22 (d, 1 H, J = 8.8 Hz), 6.87 (d, 2 H, J = 8.6 Hz), 6.81 - 6.78 (m, 5 H), 6.65 (dd, 1 H, J = 8.8 Hz, J = 2.2 Hz), 5.20 (s, 2 H), 4.27 (t, 2 H, J = 5.0 Hz), 3.44 - 3.37 (m, 4 H), 3.00 - 2.85 (m, 2 H), 1.81-1.60 (m, 5 H), 1.41 - 1.26 (m, 1 H); IR (KBr) 3350, 1470, 1250 CM<sup>-1</sup>; MS eI m/z 476 (M+); CHN calc for C<sub>28</sub>H<sub>29</sub>ClN<sub>2</sub>O<sub>3</sub> + HCl + 1.5 H<sub>2</sub>O.

**Example No. 135 3-Chloro-2-(4-hydroxy-phenyl)-1-[4-(2-azepan-1-yl-ethoxy)-benzyl]-1H-indol-5-ol (HCl)**

25 Synthesized as described for No. 134.

30 Mp = 196-198°C; <sup>1</sup>H NMR (DMSO) 10.10 (brs, 1 H), 9.86 (s, 1 H), 9.07 (s, 1 H), 7.26 (d, 2 H, J = 8.8 Hz), 7.22 (d, 1 H, J = 9.0 Hz), 6.87 (d, 2 H, J = 8.6 Hz), 6.84 - 6.78 (m, 5 H), 6.65 (dd, 1 H, J = 8.8 Hz, J = 2.2 Hz), 5.20 (s, 2 H), 4.27 (t, 2 H, J = 5.0 Hz), 3.45-3.30 (m, 4 H), 3.21-3.10 (m, 2 H), 1.82-1.76 (m, 4 H), 1.65 - 1.46 (m, 4 H); MS eI m/z 491 (M+); CHN calc for C<sub>29</sub>H<sub>31</sub>ClN<sub>2</sub>O<sub>3</sub> + 1 HCl + .37 H<sub>2</sub>O; IR (KBr) 3400, 3200, 1450, 1125

Example No. 136 3-Chloro-2-(4-hydroxy-2-methyl-phenyl)-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indol-5-ol

Synthesized as described for No. 134 except the compound was not converted into a salt.

5 Foam;  $^1\text{H}$  NMR (DMSO) 9.64 (s, 1H), 9.01 (s, 1H), 7.25 (d, 1 H,  $J$  = 8.8Hz), 7.03 (d, 1 H,  $J$  = 8.1 Hz), 6.79 (d, 1 H,  $J$  = 2.4 Hz), 6.78 - 6.65 (m, 7 H), 5.06 - 4.92 (m, 2 H), 3.94 (t, 2 H,  $J$  = 5.9 Hz), 2.62 - 2.57 (m, 2 H), 2.42 - 2.32 (m, 4 H), 1.90 (s, 3 H), 1.48- 1.40 (m, 4 H), 1.40 - 1.32 (m, 2 H); MS eI m/z 490 (M+); IR (KBr) 3430, 2900, 1450  $\text{cm}^{-1}$ ; CHN calc for 10  $\text{C}_{29}\text{H}_{31}\text{ClN}_2\text{O}_3$  + 1.0  $\text{H}_2\text{O}$ .

Synthesis of 3-ethylindole analogue No. 137

This compound was synthesized in exact analogy to the example given for 3-methylindoles, supra, using methods a and 2-8. The only difference is that the starting material used is 4'-(benzyloxy)-Butyrophenone CAS No. [26945-71-1] instead of 4'-(benzyloxy)-Propiophenone. Data for intermediates is as follows.

20 Example No. 150 5-Benzyl-2-(4-benzyl-phenyl)-3-ethyl-1H-indole

$M_p$  = 101 - 108°C; MS eI m/z 433 (M+).

Example No. 151 4-[5-Benzyl-2-(4-benzyl-phenyl)-3-ethyl-indol-1-ylmethyl-phenoxyl]-acetic acid ethyl ester

25  $M_p$  = 72 - 75°C; MS eI m/z 625 (M+).

Example No. 152 2-[4-[5-Benzyl-2-(4-benzyl-phenyl)-3-ethyl-indol-1-ylmethyl-phenoxyl]-ethanol

$M_p$  = 105 - 113 °C; MS eI m/z 583 (M+).

30

Example No. 153 Benzyl-2-(4-benzyl-phenyl)-1-[4-(2-bromo-ethoxy)-benzyl]-3-ethyl-1H-indole

$M_p$  = 140°C (decomp.); MS eI m/z 647, 645 (M+, Br present).

Example No. 154 5-Benzyl-2-(4-benzyl-phenyl)-3-ethyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole

5 Mp = 92 - 96°C; <sup>1</sup>H NMR (DMSO) 7.47 (d, 4 H, J = 7.2 Hz), 7.42 - 7.39 (m, 4 H),  
7.36 - 7.30 (m, 2 H), 7.27 (d, 2 H, J = 8.6 Hz), 7.18 (d, 1 H, J = 8.8 Hz), 7.14  
(d, 1 H, J = 2.4 Hz), 7.10 (d, 2 H, J = 8.8 Hz), 6.79 (dd, 1 H, J = 8.8 Hz, 2.2 Hz),  
6.73 (s, 4 H), 5.13 (s, 2 H), 5.11 (s, 4 H), 3.93 (t, 2 H, J = 5.9 Hz), 2.62 - 2.53  
(m, 4 H), 2.40 - 2.33 (m, 4 H), 1.49 - 1.42 (m, 4 H), 1.37 - 1.30 (m, 2 H), 1.10  
(t, 3 H, J = 7.2 Hz); MS eI m/z 650 (M+H+).

10

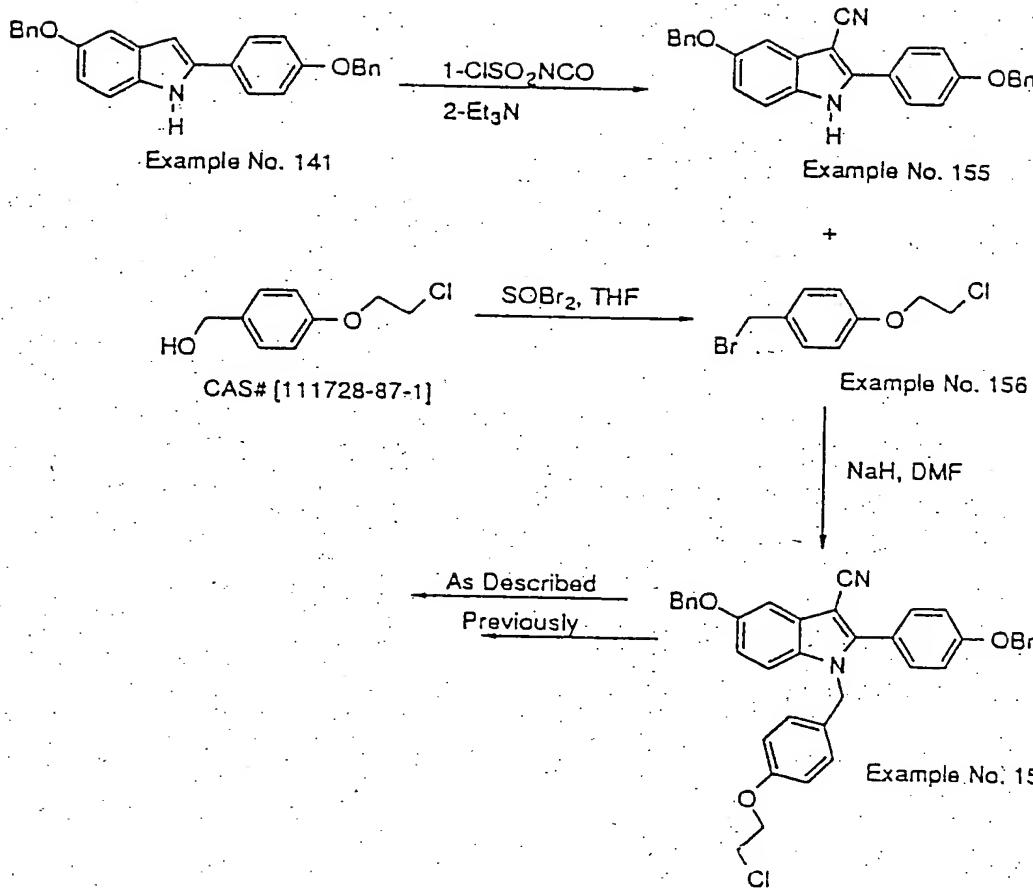
Example No. 137 2-(4-Hydroxy-phenyl)-3-ethyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indol-5-ol (HCl)

15 Mp = 160 - 164°C; <sup>1</sup>H NMR (DMSO) 9.78 (br s, 1 H), 9.69 (s, 1 H), 8.69 (s, 1 H),  
7.14 (d, 2 H, J = 8.6 Hz), 7.05 (d, 1 H, J = 8.6 Hz), 6.87 - 6.78 (m, 7 H), 6.56  
(dd, 1 H, J = 8.8 Hz, 2.4 Hz), 5.08 (s, 2 H), 4.25 (t, 2 H, J = 4.4 Hz), 3.45 - 3.38  
(m, 5 H), 3.00 - 2.86 (m, 2 H), 2.57 - 2.50 (m, 2 H), 1.83 - 1.59 (m, 5 H), 1.41 -  
1.28 (m, 1 H), 1.10 (t, 2 H, J = 7.5 Hz); IR (KBr) 3400 br, 3200 br, 2920, 1610  
cm-1; MS eI m/z 470 (M+); CHN calcd for C<sub>30</sub>H<sub>34</sub>N<sub>2</sub>O<sub>3</sub> + HCl + 1.5 H<sub>2</sub>O.

20 Scheme 15

Synthesis of 3-cyanoindole analogues

25



Example No. 155 5-Benzylxy-3-cyano-2-(4-benzyloxy-phenyl)-1H-indole

In a reaction flask 5-Benzylxy-2-(4-benzyloxy-phenyl)-1H-indole No. 141 (5.90 g, 14.6 mmol) was mixed with CH<sub>2</sub>Cl<sub>2</sub> (90 mL) was cooled down to 0°C (the starting material did not completely dissolve in CH<sub>2</sub>Cl<sub>2</sub>). While stirring vigorously, a 10 solution of chlorosulfonyl isocyanate (2.26 g, 16.0 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (25 mL) was added dropwise over a period of 45 min. The reaction was run at 0°C for 2 hrs while detected by TLC for the formation of the insoluble N-chlorosulfonylamide intermediate. After this period, Et<sub>3</sub>N (1.47 g, 14.6 mL) in CH<sub>2</sub>Cl<sub>2</sub> (25 mL) was added dropwise over 45 min at 0°C. The insoluble residue became soluble in the reaction solvent as the

Et<sub>3</sub>N addition was approaching completion. The reaction was let go for the additional 1 hr at 0°C and 2 hrs at rt. The progress of the reaction could be observed by the insoluble solid formation of the product as the reaction time went on. The solvent was stripped down and the solid residue purified by trituration with methanol to yield (4.0 g, 63.8 %). Mp = 238 - 242°C; <sup>1</sup>H NMR (DMSO) 12.31 (s, 1 H), 7.88 (d, 2 H, J = 8.8 Hz), 7.48 (d, 4 H, J = 7.25 Hz), 7.55 - 7.30 (m, 7 H), 7.23 (d, 2 H, J = 8.8 Hz), 7.14 (d, 1 H, J = 2.4 Hz), 6.97 (dd, 1 H, J = 2.2 Hz, 8.8 Hz), 5.20 (s, 2 H), 5.17 (s, 2 H); MS eI m/z 430 (M+).

10 Example No. 156 4-(2-Chloroethoxy)benzylbromide

To 4-(2-Chloroethoxy)benzylalcohol CAS No. [111728-87-1] (6.4 g, 34.31 mmol) in dioxane (100 mL) at 0°C was added slowly thionylbromide (7.13 g, 34.31 mmol). The reaction was run at 0°C after 5 min. The reaction mixture was diluted with ether (200 mL) and washed with H<sub>2</sub>O (1x30 mL) then NaHCO<sub>3</sub> (2x25 mL), and brine (30 mL). The organic extract was dried over MgSO<sub>4</sub> and concentrated. The crude product was purified by silica gel chromatography (15% EtOAc/Hex) to yield 5.0 g (58%) of the desired product Mp = 64-66°C; <sup>1</sup>H NMR (DMSO) 7.37 (d, 2 H, J = 8.8 Hz), 6.93 (d, 2 H, J = 8.8 Hz), 4.68 (s, 2 H), 4.24 (t, 2 H, J = 5.05 Hz), 3.93 (t, 2 H, J = 5.27 Hz); MS eI m/z 248 (M+).

25 Example No. 157 Benzylxy-2-(4-benzylxy-phenyl)-1-[4-(2-chloroethoxy)-benzyl]-3-cyano-1H-indole

In a reaction flask the 3-cyano indole starting material No. 155 (2.86 g, 6.64 mmol) was dissolved in DMF (25 mL) at 0°C was added NaH (191.2 mg, 8 mmol) slowly. The reaction was stirred at 0°C for 20 min. In a separate reaction flask containing 4-(2-Chloroethoxy)benzylbromide No. 156 (1.81 g, 7.28 mmol) in DMF (15 mL) at 0°C, the above prepared indole anion solution taken up by syringe was added slowly. The reaction was stirred at 0 °C for 20 min and promoted to rt for 1 h. The reaction was quenched with a few drops of H<sub>2</sub>O. The reaction mixture was partitioned between EtOAc (2x100 mL) and H<sub>2</sub>O (80 mL). The organic extract was washed with brine (80 mL), dried over MgSO<sub>4</sub>, and concentrated. The crude product was purified by trituration with ether to give the product as a white solid (2.80 g,

70.4%). Mp = 160-162°C;  $^1\text{H}$  NMR (DMSO) 7.53 - 7.28 (m, 13 H), 7.23 (m, 3 H), 6.97 (dd, 1 H, J = 2.4 Hz, 9.0 Hz), 6.86 - 6.78 (m, 4 H), 5.37 (s, 2 H), 5.18 (s, 4 H), 4.15 (t, 2 H, J = 4.8 Hz), 3.87 (t, 2 H, J = 5.3 Hz); MS eI m/z 598 (M+).

5 Example No. 's 158 and 159

Substitution of the chloro group with piperidine and hexamethyleneamine was performed analogously to the procedure outlined in method 6 using No. 157 as a starting material, *supra*.

10 Example No. 158 5-Benzyl oxy-2-(4-benzyl oxy-phenyl)-3-cyano-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole

Mp = 148 - 150 °C;  $^1\text{H}$  NMR (DMSO) 7.54 - 7.30 (m, 13 H), 7.25 - 7.18 (m, 3 H), 6.98 (dd, 1 H, J = 2.4 Hz, 9.0 Hz), 6.84 - 6.74 (m, 4 H), 5.35 (s, 2 H), 5.17 (s, 4 H), 3.94 (t, 2 H, 5.9 Hz), 2.55 (t, 2 H, 5.7 Hz), 2.35 (bs, 4 H), 1.50 - 1.40 (m, 4 H), 1.38 - 1.25 (m, 2 H); IR 3400, 2910, 2250, 1250  $\text{cm}^{-1}$ ; MS FAB 648 [M+H]<sup>+</sup>.

20 Example No. 159 5-Benzyl oxy-2-(4-benzyl oxy-phenyl)-3-cyano-1-[4-(2-azepan-1-yl-ethoxy)-benzyl]-1H-indole

$^1\text{H}$  NMR (DMSO) 8.60 (br s, 1 H), 7.60 - 7.28 (m, 12 H), 7.25 - 7.16 (m, 3 H), 6.97 (dd, 1 H, J = 2.4 Hz, 9.0 Hz), 6.88 - 6.75 (m, 4 H), 5.35 (s, 2 H), 5.17 (s, 4 H), 3.92 (t, 2 H, J = 6.2 Hz), 3.08-3.00 (m, 2 H), 2.77 (t, 2 H, J = 5.9 Hz), 2.63 (t, 4 H, J = 4.8 Hz), 1.78 - 1.68 (m, 2 H), 1.60 - 1.40 (m, 4 H); MS eI m/z 661 (M+).

Examples No. 138 and No. 139

Benzyl ethers were removed by hydrogen transfer using 1,4 cyclohexadiene and 10% Pd/C as described in method 7. Compounds were converted into their respective hydrochloride salts as described in method 8.

Example No. 138 5-Hydroxy-2-(4-Hydroxy-phenyl)-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole-3-carbonitrile (HCl)

Mp = 173 - 175°C;  $^1\text{H}$  NMR (DMSO) 10.40 (s, 1 H), 10.12 (s, 1 H), 9.40 (s, 1 H), 7.38 (m, 2 H), 7.30 (d, 1 H,  $J$  = 8.8 Hz), 7.02 - 6.90 (m, 3 H), 6.88 (s, 4 H), 6.75 (dd, 1 H,  $J$  = 2.4 Hz, 9 Hz), 5.33 (s, 2 H), 4.30 (t, 2 H,  $J$  = 4.8 Hz), 3.51 - 3.38 (m, 4 H), 2.92 (m, 2 H), 1.85 - 1.73 (m, 4 H), 1.68 - 1.59 (m, 1 H), 1.26 - 1.21 (m, 1 H); IR 3400, 2200, 1250  $\text{cm}^{-1}$ ; MS ei m/z 467 (M+); CHN calcd for  $\text{C}_{29}\text{H}_{29}\text{N}_3\text{O}_3$  + 1.0 HCl + 1.0  $\text{H}_2\text{O}$ .

10 Example No. 139 1-[4-(2-Azepan-1-yl-ethoxy)-benzyl]-5-hydroxy-2-(4-hydroxy-phenyl)-1H-indole-3-carbonitrile (HCl)

Mp = 160 - 163°C;  $^1\text{H}$  NMR (DMSO) 10.22 (s, 1 H), 10.08 (s, 1 H), 9.35 (s, 1 H), 7.40 - 7.37 (m, 2 H), 7.30 (d, 1 H, 8.8 Hz), 7.0 - 6.90 (m, 3 H), 6.87 (s, 4 H), 6.74 (dd, 1 H,  $J$  = 2.41 Hz, 9 Hz), 5.33 (s, 2 H), 4.27 (t, 2 H,  $J$  = 5.0 Hz), 3.50 - 3.30 (m, 4 H), 3.20 (m, 2 H), 1.85 - 1.70 (m, 4 H), 1.65 - 1.50 (m, 4 H); IR 3300, 2200, 1250  $\text{cm}^{-1}$ ; MS ei m/z 481 (M+); CHN calc for  $\text{C}_{30}\text{H}_{31}\text{N}_3\text{O}_3$  + 1 HCl + 1  $\text{H}_2\text{O}$ .

Esters of Indole No. 's 97 and 98

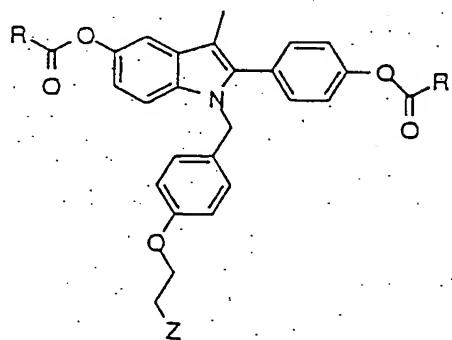


Table 9

Example No.	R	Z
No. 160	Et	
No. 161	t-Bu	
No. 162	t-Bu	

Method 9

5      Example No. 162 Di-pivalate ester of 2-(4-Hydroxy-phenyl)-3-methyl-1-[4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indol-5-ol

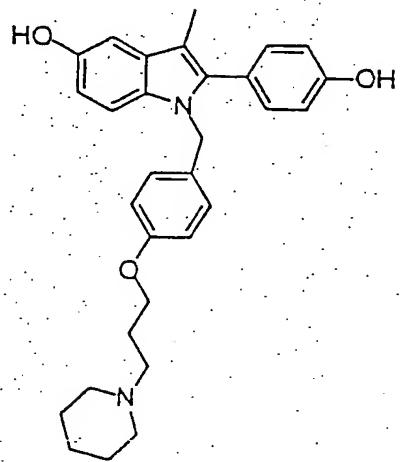
Example No. 97 free base was used as the starting material for this synthesis. No. 97 (1.0 g, 2.5 mmol) in 20 mL  $\text{CH}_2\text{Cl}_2$  was treated with diisopropylethylamine (0.7 g, 6.3 mmol) and catalytic DMAP. The reaction was cooled to 0°C and treated with pivaloyl chloride (0.7 mL, 5.6 mmol) and allowed to come to rt and stirred overnight. The reaction was worked up by diluting with  $\text{CH}_2\text{Cl}_2$  and washing with water and brine. After drying over  $\text{MgSO}_4$  the solution was concentrated and chromatographed on silica gel ( $\text{MeOH}/\text{CH}_2\text{Cl}_2$ , 1:19) to yield the desired material as an orange foam (1.08 g). This material was then taken up in 15 mL ethyl acetate and treated with 2.5 mL of a 1M  $\text{HCl}/\text{Et}_2\text{O}$  solution. Hexane was added until the solution turned cloudy. The product precipitated out as the  $\text{HCl}$  salt. This material was recrystallized from ethyl acetate/hexane to yield 0.42 g of pure No. 162:  $\text{Mp} = 182 - 185^\circ\text{C}$ ;  $\text{CHN}$  calcd for  $\text{C}_{39}\text{H}_{48}\text{N}_2\text{O}_5 + \text{HCl} + 0.25 \text{H}_2\text{O}$ .

20      Example No. 160 Di-propionate of 1-[4-(2-Azenan-1-yl-ethoxy)-benzyl]-2-(4-hydroxy-phenyl)-3-methyl-1H-indol-5-ol (HCl)  
 Compound was prepared analogously to example No. 162 except the starting material used was example No. 98 and the acylating agent used was propionyl chloride:  $\text{Mp} = 170.5 - 172^\circ\text{C}$ ;  $\text{CHN}$  calcd for  $\text{C}_{36}\text{H}_{42}\text{N}_2\text{O}_5 + \text{HCl} + 0.75 \text{H}_2\text{O}$ ; MS FAB 605 ( $\text{M}+\text{Na}$ ) $^+$ .

Example No. 161 Di-pivalate of 1-[4-(2-Azepan-1-yl-ethoxy)-benzyl-2-(4-hydroxy-phenyl)-3-methyl-1H-indol-5-ol (HCl)

Compound was prepared analogously to example No. 162 except the starting material used was example No. 98: Mp = 143 - 151°C; CHN calcd for  $C_{40}H_{50}N_2O_5$  + HCl + 0.75 H<sub>2</sub>O.

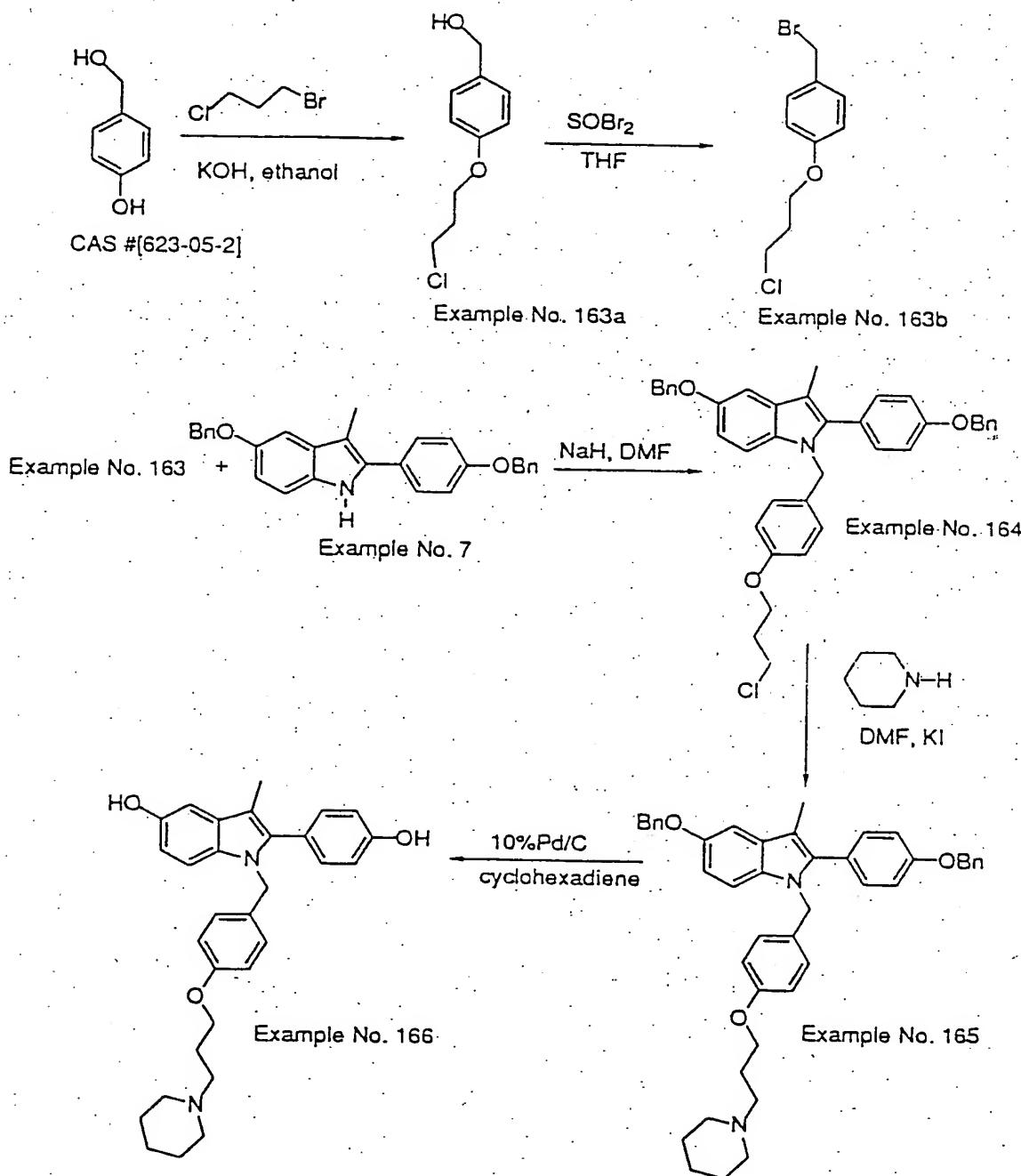
Experimental for example No. 166



10

- 90 -

Scheme 16  
Synthesis of No. 166



EXAMPLE No. 166

2-(4-Hydroxy-phenyl)-3-methyl-1-[4-[3-(piperidin-1-yl)-propoxyl]-benzyl]-1H-indol-5-ol

5 The title compound was prepared according to Scheme 16 and the steps provided below:

**Method 11**

Example No. 163a 4-(3-chloropropoxy)-benzyl alcohol

10

A solution of 4-hydroxy benzyl alcohol CAS No. [623-05-2] (10g, 80.5 mmol) in ethanol (70 mL) was treated with 1, 3 bromochloro propane (16.0g, 100 mmol) and potassium hydroxide (5.0 g, 89 mmol) was refluxed for 2 hours. The solution was cooled and filtered and then the filtrate concentrated. The concentrate was taken up in ether and washed with water, brine and dried over magnesium sulfate. The material was chromatographed on silica gel using ethyl acetate/hexanes (3:7) to yield 11.6 g of the product as a white solid: Mp = 65°C; <sup>1</sup>H NMR (DMSO) 7.21 (d, 2 H, J = 8.8 Hz), 6.88 (d, 2 H, J = 8.8 Hz), 5.03 (t, 1 H, J = 5.7 Hz), 4.40 (d, 2H, J = 5.5 Hz), 4.05 (t, 2 H, J = 6.1 Hz), 3.77 (t, 2 H, J = 6.4 Hz); MS ei m/z 200.

20

**Method 12**

Example No. 163b 4-(3-chloropropoxy)-benzyl bromide

25 A solution consisting of 4-(3-chloropropoxy)-benzyl alcohol No. 162 (10.6 g, 52.8 mmol) in dioxane (0.125 l) was cooled to 0° C and treated with a dropwise addition of thionyl bromide (12.0 g, 58.0 mmol). After 10 minutes the reaction was complete. The dioxane was diluted with ethyl ether and washed with water, brine, and then dried over MgSO<sub>4</sub>. The material was concentrated down to yield 15 g of an oil: <sup>1</sup>H NMR (DMSO) 7.36 (d, 2 H, J = 8.8 Hz), 6.92 (d, 2 H, J = 8.6 Hz), 4.68 (s, 2 H), 4.08 (t, 2 H, J = 5.9 Hz), 3.77 (t, 2 H, J = 6.4 Hz); MS (FAB) 266 (M+H<sup>+</sup>).

30

**Method 13**

**Example No. 164 5-Benzylxy-2-(4-benzylxy-phenyl)-1-[4-(3-chloro-propoxy)-benzyl]-3-methyl-1H-indole**

5 A solution consisting of 5-Benzylxy-2-(4-benzylxy-phenyl)-3-methyl-1H-indole No. 7 (6.5 g, 15.5 mmol) in DMF (60 mL) was cooled to 0°C and treated with addition of sodium hydride (0.68 g, 17.0 mmol) and stirred for 20 minutes. A solution of 4-(3-chloropropoxy)-benzyl bromide No. 163 in DMF (10 mL) was then added slowly. The reaction was allowed to come to rt and stirred for 2 hours. The reaction  
10 was poured into water and extracted with ethyl acetate. The ethyl acetate was washed with water, brine and dried over magnesium sulfate and concentrated. The concentrate was treated with methanol and 5 g of the desired product precipitated as a white solid with a melting point of 130-132°C.

15 **Method 14**

**Example No. 165 5-Benzylxy-2-(4-benzylxy-phenyl)-1-[4-(3-piperidin-1-yl-propoxy)-benzyl]-3-methyl-1H-indole**

20 A solution of 5-Benzylxy-2-(4-benzylxy-phenyl)-1-[4-(3-chloro-propoxy)-benzyl]-3-methyl-1H-indole No. 164 (3g, 5.1 mmol), potassium iodide (2.5 g, 15.3 mmol) and piperidine (3.0 mL, 30.6 mmol) were heated in DMF (30 mL) at 100°C for 18 hours. The reaction was worked up by pouring into water and extracting with ethyl acetate. The organic layer was washed with water, brine and dried over magnesium sulfate. The solution was concentrated to an oil and the product precipitated out by adding methanol. The product was obtained as a white solid: Mp = 104-106°C; <sup>1</sup>H NMR (DMSO) 7.47 (d, 4 H, J = 7.5 Hz), 7.38 (q, 4 H, J = 7.9 Hz), 7.36-7.30 (m, 1 H), 7.28 (d, 2 H, J = 8.3 Hz), 7.19 (d, 1 H, J = 8.8 Hz), 7.12-7.10 (m, 4 H), 6.80 (dd, 1 H, J = 8.8, 2.0 Hz), 6.72 (s, 4 H), 5.14 (s, 2 H), 5.13 (s, 2 H), 5.11 (s, 2 H), 3.86 (t, 2 H, J = 6.4 Hz), 2.35-2.20 (m, 6 H), 2.14 (s, 3 H), 1.78-1.75 (m, 2 H), 1.47-1.42 (m, 4 H), 1.40-1.31 (m, 2 H); MS eI m/z 650.

Method 15

Example No. 166 2-(4-Hydroxy-phenyl)-3-methyl-1-[4-[3-(piperidin-1-yl)-propoxy]-benzyl]-1H-indol-5-ol

5 A solution of 5-Benzyl-2-(4-benzyl-phenyl)-1-[4-(3-piperidin-1-yl)-propoxy]-benzyl]-3-methyl-1H-indole No. 165 (2.35 g) in tetrahydrofuran (25 mL) and ethanol (25 mL) was added to 2.3 g of 10% palladium on carbon. Cyclohexadiene (10 mL) was added and the reaction allowed to stir at room temperature for 18 hours. The catalyst was filtered through celite and the reaction mixture was concentrated and 10 chromatographed on silica gel using dichloromethane/methanol (4:1) to elute the product (0.8 g) as a white foam: Mp = 125-130°C; <sup>1</sup>H NMR 9.68 (s, 1 H), 8.70 (s, 1 H), 7.15 (d, 2 H, J = 8.6 Hz), 7.05 (d, 1 H, J = 8.8 Hz), 6.85 (d, 2 H, J = 8.6 Hz), 6.80 (d, 1 H, J = 2.4 Hz), 6.74 (d, 4 H, J = 2.6 Hz), 6.57 (dd, 1 H, J = 8.6, 2.2 Hz), 5.09 (s, 2 H), 3.88 (t, 2 H, J = 6.4 Hz), 3.60-3.15 (m, 2 H), 2.62-2.38 (m, 4 H), 2.09 (s, 3 H), 1.92-1.78 (m, 2 H), 1.55-1.43 (m, 4 H), 1.42-1.30 (m, 2 H); IR (KBr) 3400 (br), 2900, 1620, 1515 cm<sup>-1</sup>; MS ei m/z 470.

15

Synthesis of No. 167 and No. 168

20

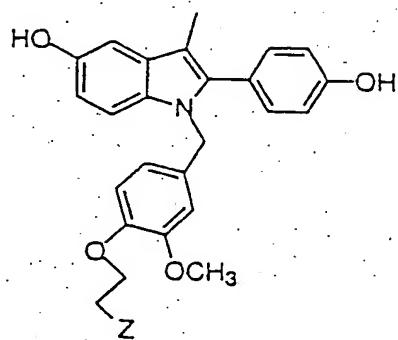
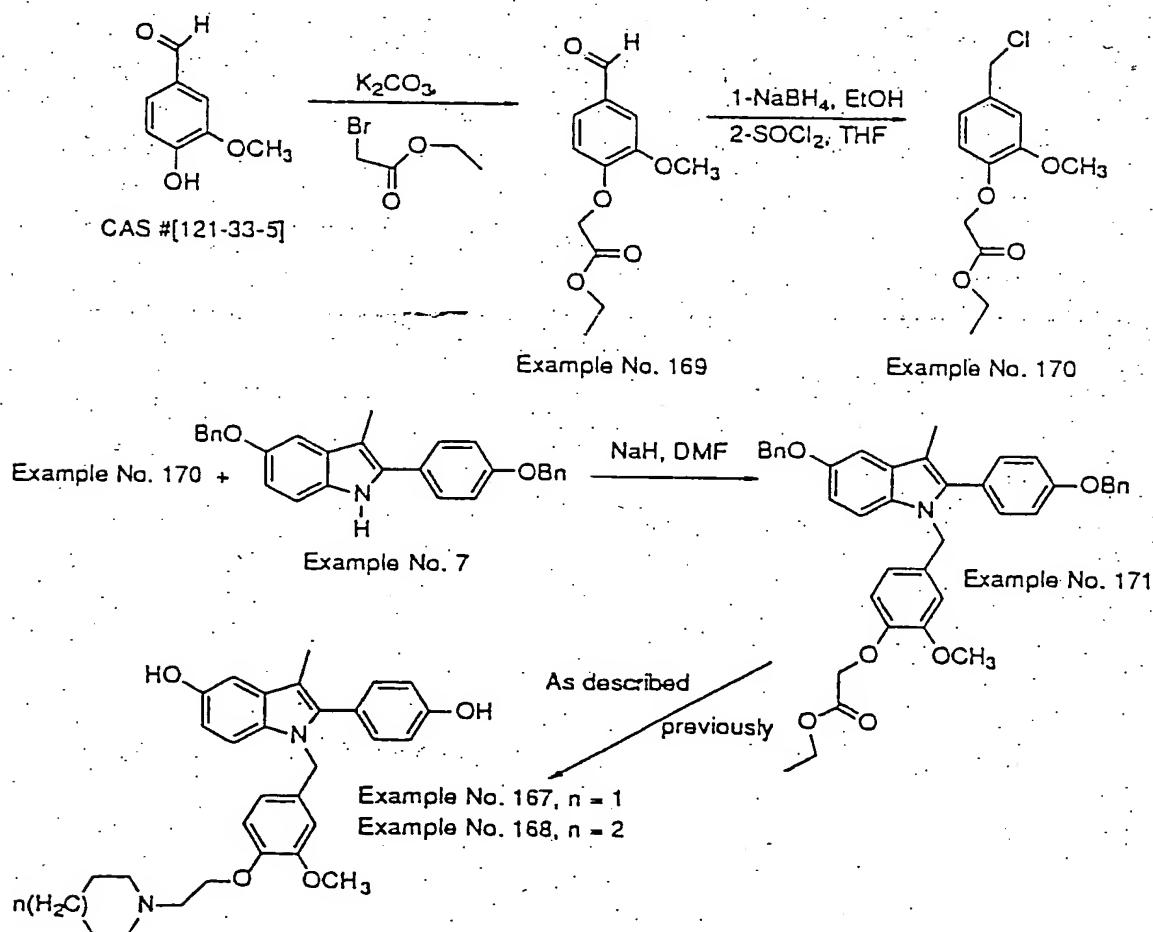


Table 10

Example No.	Z
No. 167	
No. 168	

5 Scheme 17

Synthetic scheme for examples No. 167 and No. 168



Synthesis of example No. 167

2-(4-Hydroxy-phenyl)-1-[3-methoxy-4-(2-piperidin-1-yl-ethoxy)-benzyl]-3-methyl-1H-indol-5-ol

5 Example No. 169 (4-Formyl-2-methoxy-phenoxy)-acetic acid ethyl ester

A flask containing vanillin (20g, 0.13 mol), ethyl bromoacetate (28.4g, 0.17 mol) and potassium carbonate (32.7 g, 0.24 mol) and acetone 200 mL were heated to reflux for 3 hours. The reaction was allowed to come to rt. The acetone was stripped off and the residue partitioned between water and ethyl acetate. The ethyl acetate was washed with brine and dried over magnesium sulfate. The organic layer was concentrated and the solid triturated with hexanes to yield 28.4 grams of example No. 169.

10 Mp = 56 - 59 °C;  $^1\text{H}$  NMR (DMSO) 9.83 (s, 1 H), 7.50 (dd, 1 H,  $J$  = 2.0 Hz, 8.3 Hz), 7.42 (d, 1 H,  $J$  = 1.7 Hz), 7.07 (d, 1 H,  $J$  = 8.4 Hz), 4.91 (s, 2 H), 4.16 (q, 2 H,  $J$  = 7.2 Hz), 3.84 (s, 3 H), 1.20 (t, 3 H,  $J$  = 7.1 Hz); MS ei m/z 238 (M+); CHN calcd for  $\text{C}_{12}\text{H}_{14}\text{O}_5$ .

15 Example No. 170 (4-Chloromethyl-2-methoxy-phenoxy)-acetic acid ethyl ester

20 A solution of example No. 169 (28.8g, 0.119 mol) in 600 mL of EtOH/THF(1:1) was treated with sodium borohydride (2.25 g, 0.06 mol) at 0°C and stirred for 45 minutes. The solvents were evaporated and the reaction mixture diluted with ethyl acetate and washed with 1N HCl solution. The product thus obtained (14.2 g, 0.059 mol) as an oil was dissolved in 140 mL of THF and cooled to 0°C. This solution was then treated 25 with dropwise addition of thionyl chloride (7.38g, 0.062 mol) at 0°C. After 1 hour the reaction was poured into 400 mL of water and extracted with ether. The ether layer was washed with a sodium bicarbonate solution and dried over magnesium sulfate. This was concentrated and chromatographed by silica gel chromatography using ethyl acetate/hexanes (1:9). The product was obtained as 10.5 g of a white solid. Mp = 64 - 66°C;  $^1\text{H}$  NMR (DMSO) 7.06 (d, 1 H,  $J$  = 2.0 Hz), 6.91 (dd, 1 H,  $J$  = 2.0 Hz, 2.2 Hz), 6.83 (d, 1 H,  $J$  = 2.1 Hz), 4.75 (s, 2 H), 4.70 (s, 2 H), 4.13 (q, 2 H,  $J$  = 7.2 Hz), 3.77 (s, 3 H), 1.19 (t, 3 H,  $J$  = 7.1 Hz); MS ei m/z 258 (M+); CHN calcd for  $\text{C}_{12}\text{H}_{15}\text{ClO}_4$ .

Example No. 171 2-Methoxy-4-[5-benzyl oxy-2-(4-benzyl oxy-phenyl)-3-methyl-indol-1-ylmethyl-phenoxyl]-acetic acid ethyl ester

Alkylation of the indole No. 7 was performed as described previously in Method No. 5 using example No. 170 as the electrophile.

10 Mp = 120 - 123°C;  $^1\text{H}$  NMR (DMSO) 7.48 - 7.20 (m, 13 H), 7.18 - 7.10 (m, 3 H), 6.80 (dd, 1 H, J = 2.5 Hz, 8.8 Hz), 6.64 (d, 1 H, J = 8.4 Hz), 6.52 (d, 1 H, J = 2.0 Hz), 6.24 (dd, 1 H, J = 1.9 Hz, 8.1 Hz), 5.13 (s, 4 H), 5.10 (s, 2 H), 4.61 (s, 2 H), 4.10 (q, 2 H, J = 7.0 Hz), 3.58 (s, 3 H), 2.15 (s, 3 H), 1.15 (t, 3 H, J = 7.0 Hz); MS eI m/z 641 (M+).

Example No. 172 2-[2-Methoxy-4-[5-benzyl oxy-2-(4-benzyl oxy-phenyl)-3-methyl-indol-1-ylmethyl-phenoxyl]-ethanol

Reduction of the ester No. 171 was performed as described previously in Method 4.

15 Mp = 86 - 90°C;  $^1\text{H}$  NMR (DMSO) 7.48 - 7.20 (m, 13 H), 7.18 - 7.10 (m, 3 H), 6.80 (dd, 1 H, J = 2.5 Hz, 8.8 Hz), 6.64 (d, 1 H, J = 8.4 Hz), 6.52 (d, 1 H, J = 2.0 Hz), 6.24 (dd, 1 H, J = 1.9 Hz, 8.1 Hz), 5.13 (s, 4 H), 5.10 (s, 2 H), 4.76 (t, 1 H, J = 5.5 Hz), 3.83 (t, 2 H, J = 5.1 Hz), 3.63 (q, 2 H, J = 5.3 Hz), 3.56 (s, 3 H), 2.15 (s, 3 H); MS eI m/z 599 (M+).

20 Example No. 173 5-Benzyl oxy-2-(4-benzyl oxy-phenyl)-1-[3-methoxy-4-(2-bromo-ethoxy)-benzyl-3-methyl-1H-indole

Conversion of the alcohol of example No. 172 to the bromide was performed analogously to that described in Method 5.

25 Mp = 150 - 152°C;  $^1\text{H}$  NMR (DMSO) 7.48 - 7.20 (m, 13 H), 7.18 - 7.10 (m, 3 H), 6.80 (dd, 1 H, J = 2.5 Hz, 8.8 Hz), 6.64 (d, 1 H, J = 8.4 Hz), 6.52 (d, 1 H, J = 2.0 Hz), 6.24 (dd, 1 H, J = 1.9 Hz, 8.1 Hz), 5.13 (s, 4 H), 5.10 (s, 2 H), 4.15 (t, 2 H, J = 5.3 Hz), 3.70 (t, 2 H, J = 5.7 Hz), 3.58 (s, 3 H), 2.15 (s, 3 H); MS eI m/z 661 (M+).

Example No. 174 5-Benzyl-2-(4-benzyl-phenyl)-3-methyl-1-[3-Methoxy-4-(2-piperidin-1-yl-ethoxy)-benzyl]-1H-indole

Substitution of the bromide with piperidine was performed as described previously in Method 6.

5 <sup>1</sup>H NMR (DMSO) 7.48 - 7.20 (m, 13 H), 7.18 - 7.10 (m, 3 H), 6.80 (dd, 1 H, J = 2.5 Hz, 8.8 Hz), 6.64 (d, 1 H, J = 8.4 Hz), 6.52 (d, 1 H, J = 2.0 Hz), 6.24 (dd, 1 H, J = 1.9 Hz, 8.1 Hz), 5.13 (s, 4 H), 5.10 (s, 2 H), 3.90 (t, 2 H, J = 5.7 Hz), 3.55 (s, 3 H), 2.62 - 2.50 (bs, 2 H), 2.45 - 2.30 (bs, 4 H), 2.15 (s, 3 H), 1.50 - 1.40 (m, 4 H), 1.40 - 1.35 (m, 2 H); MS FAB m/z 667 (M+H+).

10

Example No. 175 5-Benzyl-2-(4-benzyl-phenyl)-3-methyl-1-[2-Methoxy-4-(2-azepan-1-yl-ethoxy)-benzyl]-1H-indole

Reaction performed exactly as for No. 174 except hexamethyleneamine was used to displace the bromide in place of piperidine.

15 <sup>1</sup>H NMR (DMSO) 7.48 - 7.20 (m, 13 H), 7.18 - 7.10 (m, 3 H), 6.80 (dd, 1 H, J = 2.5 Hz, 8.8 Hz), 6.64 (d, 1 H, J = 8.4 Hz), 6.52 (d, 1 H, J = 2.0 Hz), 6.24 (dd, 1 H, J = 1.9 Hz, 8.1 Hz), 5.13 (s, 4 H), 5.10 (s, 2 H), 3.90 (t, 2 H, J = 5.7 Hz), 3.55 (s, 3 H), 2.85 - 2.70 (bs, 2 H), 2.70 - 2.55 (s, 4 H), 2.10 (s, 3 H), 1.60 - 1.15 (m, 8 H); MS FAB m/z 681 (M+H+)

20

Example No. 167 2-(4-Hydroxy-phenyl)-1-[3-methoxy-4-(2-piperidin-1-yl-ethoxy)-benzyl]-3-methyl-1H-indol-5-ol

Compound No. 173 was hydrogenated by transfer hydrogenation as described previously in Method 7. Compound was isolated as the hydrochloride salt by

25 dissolving in ether and treating with 1.2 equivalents of 1N ether/HCl solution (this is a variation of method 8).

Mp = 123 - 127 °C; <sup>1</sup>H NMR (DMSO) 10.20 (bs, 1 H), 9.72 (s, 1 H), 8.71 (s, 1 H), 7.17 (d, 2 H, J = 8.6 Hz), 7.11 (d, 1 H, J = 8.8 Hz), 6.87 (d, 2 H, J = 8.6 Hz), 6.79 (m, 2 H), 6.57 (dd, 1 H, J = 2.4 Hz, 8.8 Hz), 6.55 (d, 1 H, J = 1.7 Hz), 6.33

30 (dd, 1 H, J = 1.7 Hz, 8.1 Hz), 5.11 (s, 2 H), 4.23 (t, 2 H, J = 4.8 Hz), 3.60 (s, 3 H), 3.45 (m, 2 H), 3.35 (m, 2 H), 2.95 (m, 2 H), 2.10 (s, 3 H), 1.70 (m, 5 H), 1.35

(m, 1 H); IR 3500, 1500, 1275  $\text{cm}^{-1}$ ; MS (+) FAB m/z 487 ( $\text{M}+\text{H}$ )<sup>+</sup>; CHN calcd for  $\text{C}_{30}\text{H}_{34}\text{N}_2\text{O}_4$  + 1 HCl + 1.0  $\text{H}_2\text{O}$ .

5 Example No. 168 2-(4-Hydroxy-phenyl)-1-[3-methoxy-4-(2-azepan-1-yl-ethoxy)-benzyl]-3-methyl-1H-indol-5-ol

Prepared in the same way as that described for example No. 167.

10 Mp = 142 - 146 °C; <sup>1</sup>H NMR (DMSO) 10.36 (s, 1 H), 9.72 (s, 1 H), 8.71 (s, 1 H), 7.18 (d, 2 H, J = 8.3 Hz), 7.11 (d, 1 H, J = 8.6 Hz), 6.87 (d, 2 H, J = 8.3 Hz), 6.82 (d, 1 H, J = 8.1 Hz), 6.79 (d, 1 H, J = 2.2 Hz), 6.57 (dd, 1 H, J = 2.2 Hz, 8.6 Hz), 6.55 (d, 1 H, J = 1.8 Hz), 6.33 (dd, 1 H, J = 1.5 Hz, 8.1 Hz), 5.11 (s, 2 H), 4.24 (t, 2 H, J = 4.6 Hz), 3.60 (s, 3 H), 3.40 (m, 4 H), 3.20 (m, 2 H), 2.10 (s, 3 H); 1.75 (m, 4 H), 1.55 (m, 4 H); IR (KBr) 3300, 1500, 1270, 1200  $\text{cm}^{-1}$ ; MS (+) FAB m/z 501 ( $\text{M}+\text{H}$ )<sup>+</sup>; CHN calcd for  $\text{C}_{31}\text{H}_{36}\text{N}_2\text{O}_4$  + 1.0 HCl + 0.12  $\text{CH}_3\text{OH}$ .

15 Biological Data

Method 16

In vitro estrogen receptor binding assay

20 Receptor preparation

CHO cells overexpressing the estrogen receptor were grown in 150  $\text{mm}^2$  dishes in DMEM + 10% dextran coated charcoal, stripped fetal bovine serum. The plates were washed twice with PBS and once with 10mM Tris-HCl, pH 7.4, 1mM EDTA. Cells 25 were harvested by scraping the surface and then the cell suspension was placed on ice. Cells were disrupted with a hand-held motorized tissue grinder using two, 10-second bursts. The crude preparation was centrifuged at 12,000g for 20 minutes followed by a 60 minute spin at 100,000g to produce a ribosome free cytosol. The cytosol was then frozen and stored at -80°C. Protein concentration of the cytosol was estimated using 30 the BCA assay with reference standard protein.

### Binding assay conditions

The competition assay was performed in a 96-well plate (polystyrene\*) which binds <2.0% of the total input [<sup>3</sup>H]-17 $\beta$ -estradiol and each data point was gathered in 5 triplicate. 100uG/100uL of the receptor preparation was aliquoted per well. A saturating dose of 2.5 nM [<sup>3</sup>H]17 $\beta$ -estradiol + competitor (or buffer) in a 50 uL volume was added in the preliminary competition when 100x and 500x competitor were evaluated, only 0.8 nM [<sup>3</sup>H] 17 $\beta$ -estradiol was used. The plate was incubated at room temperature for 2.5 h. At the end of this incubation period 150 uL of ice-cold dextran coated charcoal (5% activated charcoal coated with 0.05% 69K dextran) was added to each well and the plate was immediately centrifuged at 99g for 5 minutes at 4°C. 200 uL of the supernatant solution was then removed for scintillation counting. Samples were counted to 2% or 10 minutes, whichever occurs first. Because polystyrene absorbs a small amount of [<sup>3</sup>H]17 $\beta$ -estradiol, wells containing radioactivity and cytosol, but not processed with charcoal were included to quantitate amounts of available isotope. Also, wells containing radioactivity but no cytosol were processed with charcoal to estimate unremovable DPM of [<sup>3</sup>H] 17 $\beta$ -estradiol. Corning No. 15. 25880-96, 96-well plates were used because they have proven to bind the least amount of estradiol.

20

### Analysis of results

Counts per minute (CPM) of radioactivity were automatically converted to disintegrated per minute (DPM) by the Beckman LS 7500 Scintillation Counter using a 25 set of quenched standards to generate a H No. for each sample. To calculate the % of estradiol binding in the presence of 100 or fold 500 fold competitor the following formula was applied:

30 
$$((DPM_{sample} - DPM_{not\ removed\ by\ charcoal}) / (DPM_{estradiol} - DPM_{not\ removed\ by\ charcoal})) \times 100\% = \% \text{ of estradiol binding}$$

For the generation of IC<sub>50</sub> curves, % binding is plotted vs compound. IC<sub>50</sub>'s are generated for compounds that show >30% competition at 500x competitor

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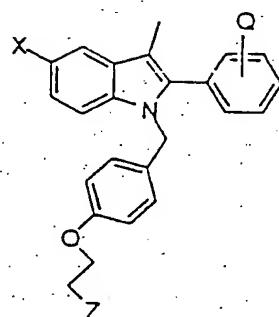
concentration. For a description of these methods, see Hulme, E.C., ed. 1992. Receptor-Ligand Interactions: A Practical Approach. IRL Press, New York.(see especially chapter 8).

5

Table 11

Estrogen Receptor Binding

10



Example No.	X	Q	Z	Receptor Binding IC50's uM
No. 85	H	H	N	0.45
No. 86	H	4'-OH	N	0.12
No. 87	OH	H	N	0.030
No. 88	OMe	4'-OH	N	0.35
No. 89	OH	4'-OMe	N	0.30
No. 90	OMe	4'-OMe	N	0.60
No. 91	OMe	4'-OMe	N	0.52
No. 92	OH	4'-OEt	N	0.062

Table 11 (Cont'd)

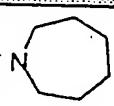
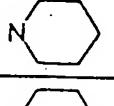
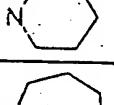
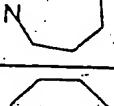
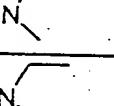
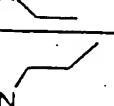
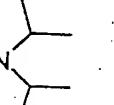
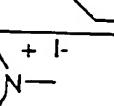
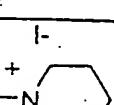
Example No.	X	Q	Z	Receptor Binding IC <sub>50</sub> $\mu$ M
No. 93	OH	4'-OEt		0.090
No. 94	F	4'-OH		0.20
No. 97	OH	4'-OH		0.060
No. 98	OH	4'-OH		0.050
No. 99	OH	4'-OH		0.03
No. 100	OH	4'-OH		0.06
No. 101	OH	4'-OH		0.04
No. 102	OH	4'-OH		0.08
No. 103	OH	4'-OH		0.2
No. 104	OH	4'-OH		0.1
No. 105	OH	4'-OH		0.028
No. 106	OH	4'-OH		0.1
No. 107	OH	4'-OH		0.06

Table 11 (Cont'd)

Example No.	X	Q	Z	Receptor Binding IC <sub>50</sub> nM
No. 108	OH	4'-OH		0.02
No. 109	OH	4'-OH		0.17
No. 110	OH	4'-OH		0.037
No. 111	OH	4'-OH		0.15
No. 112	OH	4'-OH		0.07
No. 113	OH	4'-OH		0.047
No. 114	OH	4'-OH		0.001
No. 115	OH	4'-OH		0.15
No. 116	OH	4'-Fl		0.04
No. 117	OH	4'-Fl		0.10
No. 118	OH	3'-OMe,4'-OH		N/A
No. 119	OH	3',4'-OCH <sub>2</sub> O-		0.070

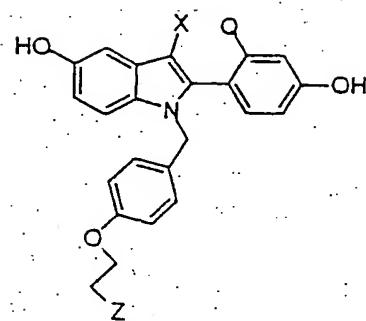
Table 11 (Cont'd)

Example No	X	Q	Z	Receptor Binding IC <sub>50</sub> 's μM
No. 120	OH	4'-O-iPr		0.10
No. 121	OH	4'-O-iPr		0.080
No. 122	OH	4'-O-Cp		0.080
No. 123	OH	4'-CF <sub>3</sub>		0.17
No. 124	OH	4'-CH <sub>3</sub>		0.11
No. 125	OH	4'-Cl		0.11
No. 126	OH	2',4',-Dimethoxy		N/A
No. 127	OH	3'-OH		0.019
No. 128	OH	3'-OH		0.009
No. 129	OH	4'-OH,3'-Fl		0.0055
No. 130	OH	4'-OH, 3'-Fl		0.013
No. 131	OH	3'-OMe		0.12
No. 132	OH	4'-OCF <sub>3</sub>		0.05

Table 12

Estrogen Receptor Binding

5

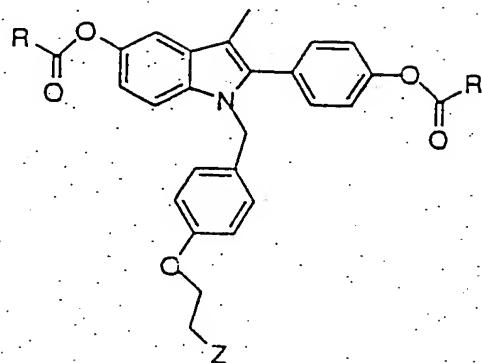


Example No.	X	Q	Z	Receptor Binding IC <sub>50</sub> nM
No. 133	Cl	H	N	0.004
No. 134	Cl	H	N	0.024
No. 135	Cl	H	N	0.029
No. 136	Cl	CH <sub>3</sub>	N	0.013
No. 137	Et	H	N	0.15
No. 138	CN	H	N	0.011
No. 139	CN	H	N	0.023

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Table 13

Estrogen Receptor Binding

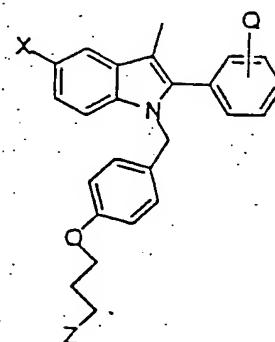


Example No.	R	Z	Receptor Binding IC50's $\mu$ M
No. 160	Et	N	N/A
No. 161	t-Bu	N	N/A
No. 162	t-Bu	N	Does not Bind

10

Table 14

Estrogen Receptor Binding



15

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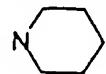
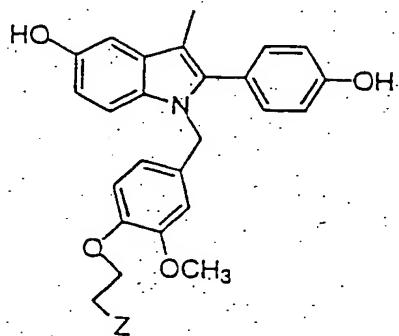
Example No.	X	Q	Z	Receptor Binding IC <sub>50</sub> 's $\mu$ M
No. 166	OH	4'-OH		0.099

Table 15

5 Estrogen Receptor Binding



10

Example No.	Z	Receptor Binding IC <sub>50</sub> 's $\mu$ M
No. 167		0.08
No. 168		0.057

Method 17

15 Ishikawa Cell Alkaline Phosphatase Assay

Cell Maintenance and Treatment:

20 Ishikawa cells were maintained in DMEM/F12 (50%:50%) containing phenol red + 10% fetal bovine serum and the medium was supplemented with 2 mM Glutamax, 1% Pen/Strap and 1 mM sodium pyruvate. Five days prior to the beginning

of each experiment (treatment of cells) the medium was changed to phenol red-free DMEM/F12 + 10% dextran coated charcoal stripped serum. On the day before treatment, cells were harvested using 0.5% trypsin/EDTA and plated at a density of  $5 \times 10^4$  cells/well in 96-well tissue culture plates. Test compounds were dosed at  $10^{-6}$ ,  $10^{-7}$  and  $10^{-8}$  M in addition to  $10^{-6}$  M (compound) +  $10^{-9}$  M 17 $\beta$ -estradiol to evaluate the ability of the compounds to function as antiestrogens. Cells were treated for 48 h prior to assay. Each 96-well plate contained a 17 $\beta$ -estradiol control. Sample population for at each dose was n=8.

10 Alkaline Phosphatase Assay:

At the end of 48h the media is aspirated and cells are washed three times with phosphate buffered saline (PBS). 50 $\mu$ L of lysis buffer (0.1 M Tris-HCl, pH 9.8, 0.2% Triton X-100) is added to each well. Plates are placed at -80°C for a minimum of 15 minutes. Plates are thawed at 37°C followed by the addition of 150 $\mu$ L of 0.1 M Tris-HCl, pH 9.8, containing 4 mM para-nitrophenylphosphate (pNPP) to each well (final concentration, 3 mM pNPP).

20 Absorbance and slope calculations were made using the KineticCalc Application program (Bio-Tek Instruments, Inc., Winooski, VT). Results are expressed as the mean +/- S.D. of the rate of enzyme reaction (slope) averaged over the linear portion of the kinetic reaction curve (optical density readings every 5 minutes for 30 minutes absorbance reading). Results for compounds are summarized as percent of response related to 1 nM 17 $\beta$ -estradiol.

25 Various compounds were assayed for estrogenic activity by the alkaline phosphatase method and corresponding ED50 values (95% C.I.) were calculated. The four listed in the following were used as reference standards:

30	17 $\beta$ -estradiol	0.03 nM
	17 $\alpha$ -estradiol	1.42 nM
	estriol	0.13 nM
	estrone	0.36 nM

A description of these methods is described by Holinka, C.F., Hata, H., Kuramoto, H. and Gurge, E. (1986) Effects of steroid hormones and antisteroids on alkaline phosphatase activity in human endometrial cancer cells (Ishikawa Line).  
5 Cancer Research, 46:2771-2774, and by Littlefield, B.A., Gurge, E., Markiewicz, L., McKinley, B. and Hochberg, R.B. (1990) A simple and sensitive microtiter plate estrogen bioassay based on stimulation alkaline phosphatase in Ishikawa cells; Estrogen action of D5 adrenal steroids. Endocrinology, 6:2757-2762.

10 Ishikawa Alkaline Phosphatase Assay

	<u>Compound</u>	<u>% Activation</u>
	17 $\beta$ -estradiol	100% activity
	tamoxifen	0% activity (45% with 1 nM 17 $\beta$ -estradiol)
15	raloxifene	5% activity (5% with 1 nM 17 $\beta$ -estradiol)
	Example No. 98	1% activity (1% with 1 nM 17 $\beta$ -estradiol)

Method No. 18

20 2X VIT ERE Infection Assay

Cell Maintenance and Treatment

Chinese Hamster Ovary cells (CHO) which had been stably transfected with the 25 human estrogen receptor were maintained in DMEM + 10% fetal bovine serum (FBS). 48h prior to treatment the growth medium was replaced with DMEM lacking phenol red + 10% dextran coated charcoal stripped FBS (treatment medium). Cells were plated at a density of 5000 cells/well in 96-well plates containing 200  $\mu$ L of medium/well.

30 Calcium Phosphate Transfection

Reporter DNA (Promega plasmid pGL2 containing two tandem copies of the vitellogenin ERE in front of the minimal thymidine kinase promoter driving the

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luciferase gene) was combined with the B-galactosidase expression plasmid pCH110 (Pharmacia) and carrier DNA (pTZ18U) in the following ratio:

10uG of reporter DNA  
5 uG of pCH110DNA  
5 uG of pTZ18U  
20 uG of DNA/1 mL of transfection solution

10 The DNA (20uG) was dissolved in 500 uL of 250 mM sterile  $\text{CaCl}_2$  and added dropwise to 500 uL of 2 X HeBS (0.28 M NaCl, 50 mM HEPES, 1.5 mM  $\text{Na}_2\text{HPO}_4$ , pH 7.05) and incubated at room temperature for 20 minutes. 20 uL of this mixture was added to each well of cells and remained on the cells for 16 h. At the end of this incubation the precipitate was removed, the cells were washed with media, fresh treatment media was replaced and the cells were treated with either vehicle, 1 nM 17 $\beta$ -estradiol, 1uM compound or 1 uM compound + 1 nM 17 $\beta$ -estradiol (tests for estrogen antagonism). Each treatment condition was performed on 8 wells (n=8) which were incubated for 24 h prior to the luciferase assay.

#### Luciferase Assay

20 After 24h exposure to compounds, the media was removed and each well washed with 2 X with 125 uL of PBS lacking  $\text{Mg}^{++}$  and  $\text{Ca}^{++}$ . After removing the PBS, 25 uL of Promega lysis buffer was added to each well and allowed to stand at room temperature for 15 min, followed by 15 min at -80°C and 15 min at 37°C. 20 uL of lysate was transferred to an opaque 96 well plate for luciferase activity evaluation and the remaining lysate (5 uL) was used for the B-galactosidase activity evaluation (normalize transfection). The luciferan substrate (Promega) was added in 100 uL aliquots to each well automatically by the luminometer and the light produced (relative light units) was read 10 seconds after addition.

#### Infection Luciferase Assay (Standards)

<u>Compound</u>	<u>% Activation</u>
-----------------	---------------------

17 $\beta$ -estradiol	100% activity
estriol	38% activity
tamoxifen	0% activity (10% with 1 nM 17 $\beta$ -estradiol)
raloxifene	0% activity (0% with 1 nM 17 $\beta$ -estradiol)

5

### B-Galactosidase Assay

To the remaining 5  $\mu$ L of lysate 45  $\mu$ L of PBS was added. Then 50  $\mu$ L of Promega B-galactosidase 2X assay buffer was added, mixed well and incubated at 10 37°C for 1 hour. A plate containing a standard curve (0.1 to 1.5 milliunits in triplicate) was set up for each experimental run. The plates were analyzed on a Molecular Devices spectrophotometric plate reader at 410 nm. The optical densities for the unknown were converted to milliunits of activity by mathematical extrapolation from the standard curve.

15

### Analysis of Results

The luciferase data was generated as relative light units (RLUs) accumulated during a 10 second measurement and automatically transferred to a JMP (SAS Inc) file 20 where background RLUs were subtracted. The B-galactosidase values were automatically imported into the file and these values were divided into the RLUs to normalize the data. The mean and standard deviations were determined from a n=8 for each treatment. Compounds activity was compared to 17 $\beta$ -estradiol for each plate. Percentage of activity as compared to 17 $\beta$ -estradiol was calculated using the formula 25  $\% = ((\text{Estradiol-control}) / (\text{compound value})) \times 100$ . These techniques are described by Tzukerman, M.T., Esty, A., Santiso-Mere, D., Danielian, P., Parker, M.G., Stein, R.B., Pike, J.W. and McDonnel, D.P. (1994). Human estrogen receptor transactivational capacity was determined by both cellular and promoter context and mediated by two functionally distinct intramolecular regions (see Molecular 30 Endocrinology, 8:21-30).

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Table 16  
 Infection Luciferase Activity

5

Example No.	1 uM	1 uM + 17 $\beta$ estradiol
No. 85	-2	43
No. 86	-5	2
No. 87	0	0
No. 88	4	44
No. 89	16	18
No. 90	3	58
No. 91	-3	56
No. 92	-4	-2
No. 93	-3	-2
No. 94	-5	15
No. 95	-4	-4
No. 96	12	8
No. 97	-4	-5
No. 98	5	5
No. 99	5	6
No. 100	9	10
No. 101	14	9
No. 102	9	10
No. 103	13	10
No. 104	7	7
No. 105	5	5
No. 106	10	81
No. 107	-1	54
No. 108	11	10
No. 109	6	5
No. 110	8	10
No. 111	25	23
No. 112	10	10
No. 113	14	16
No. 114	1	-1
No. 115	11	10
No. 116	-1	1
No. 117	0	1
No. 118	N/A	N/A
No. 119	-1	-1
No. 120	-1	1
No. 121	0	1
No. 122	1	5
No. 123	-1	1

Table 16 (Cont'd)  
 Infection Luciferase Activity

Example No.	I uM	1 uM + 17 $\beta$ estradiol
No. 124	-2	-2
No. 125	-3	-2
No. 126	-1	0
No. 127	-3	-4
No. 132	-5	-2
No. 133	7	9
No. 134	-9	5
No. 135	7	3
No. 136	16	10
No. 137	6	8
No. 138	-2	-1
No. 139	-12	-13
No. 160	N/A	N/A
No. 161	N/A	N/A
No. 162	-14	-13
No. 166	25	23
No. 167	4	10
No. 168	3	7

5

Method No. 19

Rat Uterotrophic/Antiuterotrophic Bioassay

The estrogenic and antiestrogenic properties of the compounds were determined  
 10 in an immature rat uterotrophic assay (4 day) that (as described previously by L.J.Black  
 and R.L.Goode, Life Sciences, 26, 1453 (1980)). Immature Sprague-Dawley rats  
 (female, 18 days old) were tested in groups of six. The animals were treated by daily  
 ip injection with 10 uG compound, 100 uG compound, (100 uG compound + 1 uG  
 17 $\beta$ -estradiol) to check antiestrogenicity, and 1 uG 17 $\beta$ -estradiol, with 50%  
 15 DMSO/50% saline as the injection vehicle. On day 4 the animals were sacrificed by  
 CO<sub>2</sub> asphyxiation and their uteri were removed and stripped of excess lipid, any fluid  
 removed and the wet weight determined. A small section of one horn was submitted  
 for histology and the remainder used to isolate total RNA in order to evaluate  
 complement component 3 gene expression.

Table 17  
 3 Day Rat Immature Uterine Assay

Example No.	Uterine wt mg	Uterine wt mg	Uterine wt mg	Uterine wt mg
	100 $\mu$ G cmpd	100 $\mu$ G cmpd + 1 $\mu$ G 17 $\beta$ -estradiol	1 $\mu$ G 17 $\beta$ -estradiol	Vehicle
Tamoxifen	71.4 mg	N/A	98.2 mg	42.7 mg
No. 85	41.1 mg	92.4 mg	94.4 mg	26.6 mg
No. 94	28.1 mg	93.7 mg	88.5 mg	22.3 mg
No. 97	27.4 mg	24.3 mg	63.2 mg	30.7 mg
No. 98	29.4 mg	27.9 mg	94.1 mg	35.9 mg
No. 100	59.9 mg	68.7 mg	91.9 mg	23.4 mg
No. 101	65.1 mg	71.0 mg	113.7 mg	27.7 mg
No. 122	46.7 mg	38.7 mg	103.4 mg	30.3 mg
No. 123	39.2 mg	61.4 mg	94.4 mg	26.1 mg
No. 138	28.4 mg	37.9 mg	93.9 mg	24.6 mg
No. 139	30.4 mg	45.0 mg	82.1 mg	20.5 mg
No. 168	43.2 mg	81.7 mg	98.9 mg	25.5 mg

Method No. 20

10 6-Week Ovariectomized Rat Model

Female Sprague Dawley CD rats, ovx or sham ovx, were obtained 1 day after surgery from Taconic Farm (weight range 240 - 275 g). They were housed 3 or 4 rats/cage in a room on a 14/10 (light/dark) schedule and provided with food (Purina 15 500 rat chow) and water ad libitum. Treatment for all studies began 1 day after the animals arrival and dosed 5 or 7 days per week as indicated for 6 weeks. A group of age matched sham operated rats not receiving any treatment served as an intact, estrogen replete control group for each study. All treatments were prepared in 1% tween 80 in normal saline at defined concentrations so that the treatment volume was 20 0.1mL/100g body weight. 17-beta estradiol was dissolved in corn oil (20  $\mu$ G/mL) and delivered subcutaneously, 0.1 mL/rat. All dosages were adjusted at three week intervals according to group mean body weight measurements.

5       Five weeks after the initiation of treatment and one week prior to the termination of the study, each rat was evaluated for bone mineral density (BMD). The BMD's of the proximal tibiae (PT) and fourth lumbar vertabrate (L4) were measured in anesthetized rats using a dual energy X-ray absorptiometer (Eclipse XR-26, Norland Corp. Ft. Atkins, WI). The dual energy X-ray absorptiometer (DXA) measurements for each rat were performed as follows: Fifteen minutes prior to DXA measurements, the rat was anesthetized with an intraperitoneal injection of 100 mg/kg ketamine (Bristol Laboratories, Syracuse, NY) and 0.75 mg/kg acepromazine (Aveco, Ft. Dodge, IA).

10      The rat was placed on an acrylic table under the DXA scanner perpendicular to its path; the limbs were extended and secured with paper tape to the surface of the table. A preliminary scan was performed at a scan speed of 50 mm/second with a scan resolution of 1.5 mm X 1.5 mm to determine the region of interest in PT and L4. Small subject software was employed at a scan speed of 10mm/second with resolution of 0.5 mm X 0.5 mm for final BMD measurements. The software allows the operator to define a 1.5 cm wide area to cover the total length of L4. The BMDs for respective sites were computed by the software as a function of the attenuation of the dual beam (46.8 KeV and 80 KeV) X-ray generated by the source underneath the subject and the detector travelling along the defined area above the subject. The data for BMD values (expressed in g/cm<sup>2</sup>) and individual scans were stored for statistical analysis.

15      One week after BMD evaluation the rats were sacrificed by carbon dioxide suffocation and blood collected for cholesterol determination. The uteri were removed and the weights taken. Total cholesterol is determined using a Boehringer-Mannheim Hitachi 911 clinical analyzer using the Cholesterol/HP kit. Statistics were compared using one-way analysis of variance with Dunnet's test.

20

25

Table 18

6-Week Ovariectomized Rat Study Of Example No. 98

5

Treatment	BMD (mg/cm <sup>2</sup> ) <sup>a,b</sup>		Δ Body Weight (g) <sup>c,d</sup>	Uterine Weight (mg) <sup>a,c</sup>	Cholesterol (mg/dl) <sup>a,c</sup>
	Proximal Tibia	L <sub>4</sub>			
Study <sup>a</sup>					
Sham (Intact)	0.211 <sup>**</sup> ±0.003	0.183 <sup>**</sup> ±0.003	43.0 <sup>*</sup> ±6.0	426.4 <sup>**</sup> ±25.0	71.6 <sup>**</sup> ±5.0
Vehicle (Ovx)	0.189 ±0.004	0.169 ±0.004	62.7 ±8.2	118.2 <sup>**</sup> ±7.8	87.2 <sup>**</sup> ±3.0
Example No. 98					
0.3mg/kg, p.o.	0.210 <sup>**</sup> ±0.003	0.173 ±0.003	46.8 ±6.6	149.3 ±4.4	59.0 <sup>**</sup> ±2.2
Raloxifene 3mg/kg, p.o.	0.207 <sup>**</sup> ±0.006	0.170 ±0.003	25.3 <sup>**</sup> ±5.4	191.6 <sup>**</sup> ±9.3	55.0 <sup>**</sup> ±2.4
17 $\beta$ -Estradiol 2 $\mu$ g/rat, s.c.	0.224 <sup>**</sup> ±0.004	0.169 ±0.004	33.1 <sup>**</sup> ±4.9	426.0 <sup>**</sup> ±18.4	95.5 <sup>**</sup> ±3.9

<sup>a</sup> Mean ± SEM

<sup>b</sup> Following 5 weeks of treatment

<sup>c</sup> Following 6 weeks of treatment

<sup>d</sup> Daily treatment x7 days/week x6 weeks

<sup>\*</sup>p < 0.05 vs corresponding Vehicle value

<sup>\*\*</sup>p < 0.01 vs corresponding Vehicle value

10